

About the Author



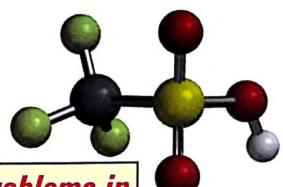
Mahendra Singh Chouhan (MSC Sir) is a renowned name in the realm of Organic Chemistry. Through a Chemical Engineer from Mumbai University, his great passion for the subject led him to impart guidance to IIT-JEE aspirants on a regular basis. His in depth knowledge and vast experience has helped innumerable students to achieve their dream of excelling at IIT, JEE and other such tough challenges.

He has launched a website to extend the benefits of his expertise beyond the geographical barriers to all those who dare to dream and seek-www.iitjeeorganic.com.

The website provides expert guidance in all the areas of the subject in a most skillful manner. There are quizzes, challenging questions, notes, e-books and videos etc. This website is a complete guide in itself for organic chemistry and has been designed for IIT-JEE aspirants, keeping in mind the various syllabi and CBSE.

Highly recommended for the high flyers.





Advanced Problems in

ORGANIC CHEMISTRY

for



by: M.S. Chouhan

Director

Vibrant Academy, Kota



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A few words to the JEE Aspirants

Dear JEE aspirants,

I hope that this collection of problems will surely help you during your preparation for JEE. In this book, each chapter consists of two levels:

Level 1 - includes the problems having only one option correct. These problems are based on different facts and their twists.

Level 2 - includes unique approach which may be used to solve the problems altogether different from the prevailing trend followed by JEE. These approaches will undoubtedly help you in the quick revision of the key facts and their applications.

I wish all of you a grand success in the ensuing Joint Entrance Examination. Your valuable suggestions and constructive criticism for the betterment of the book are welcome.

M.S. Chouhan

Preface

It is a matter of great pleasure for me to present the eleventh edition of "Advanced Problems in Organic Chemistry for JEE" before JEE aspirants. During my teaching experience, I felt that the facts may be made more and more clear to the students through problematic approach. Although an ocean of material in Organic Chemistry is available with the students, yet the approach to design the problems has been changed in recent years and if one tries to swim in this ocean, it will be a very difficult task. To make the students more familiar with trends and tricks how to solve problems, the present problem book has been presented. In the current scenario of stiff competition especially for JEE, one must be clear that almost all the sincere applicants are well equipped with the facts of subject, yet the winner is one who knows how to use these equipments with accuracy and efficiency. As an experienced teacher, I would like to suggest students three golden rules to score high in Organic Chemistry:

- 1. Don't get behind
- 2. Work out a number of problems of different types
- 3. Revise through short notes / learning chart.

I hope that the present book will cater to the needs of JEE aspirants & as a matter of fact, they will enjoy the present venture and I would feel rewarded if this book is found helpful to the students and teachers in real terms. All attempts have been made to make the book error free however a few misprints may inadvertently creep.

I acknowledge the blessing and support of my mother Smt. Raj Kanwar, father Shri B.S. Chouhan, brother Dr. V.S. Chouhan, my wife and daughter. They inspired me all the time during the preparation of this book.

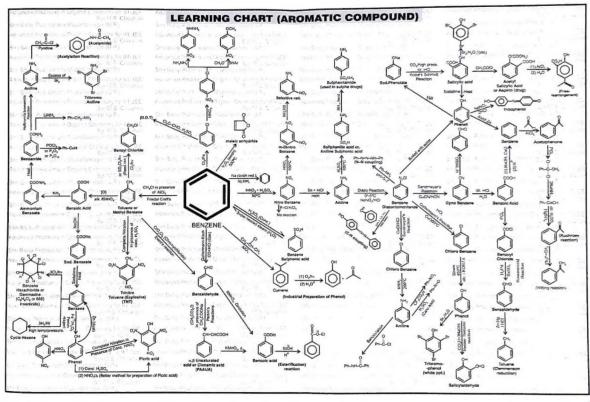
The support and valuable suggestions from my colleagues especially Mr. N. Avasthi, Mr. V. K. jaiswal, Mr. Nitin Jain, Mr. N.K. Sethia, Mr. Vikash Gupta, Mr. Pankaj Joshi, Dr. S. Kothari, Mr. Vineet Khatri, Mr. Ashish Mishra, Mr. Manish Arora, Mr. Govind Khandelwal, Mr. Rahul Pareek, Mr. Rahul Malav, Mr. Divyesh Tiwari, Mr. Omkar Kelapure, Mr. Kishore Kilani, Mr. Mayank Pareek, Mr. Gurpreet Singh, Mr. Yogesh Jain, Madam Anjana Kamal, Mr. Aneet Choudhary, Mr. Shaliwahan Singh Rathore, Mr. Akshay Chaudhary, Mr. Hanuman Sahay, Mrs. Neha Joshi, Mrs. Neetu Jha, Mr. Kamlesh Gupta and Mr. Kumud Ranjan are highly acknowledged. I also pay my sincere thanks to all the esteemed members of M/s Shri Balaji Publications in bringing out this book in such a nice form.

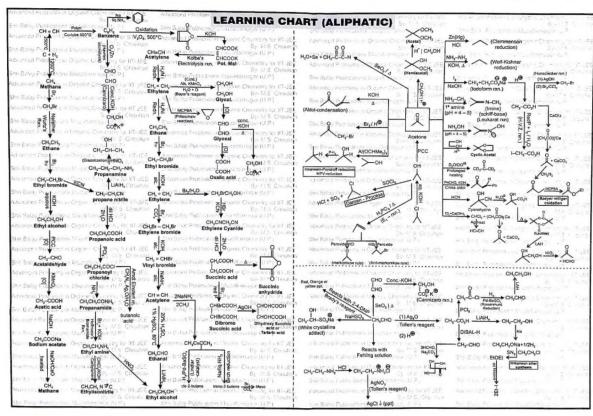
In the last, constructive criticism and valuable suggestions from the readers are most welcome to make the book more useful.

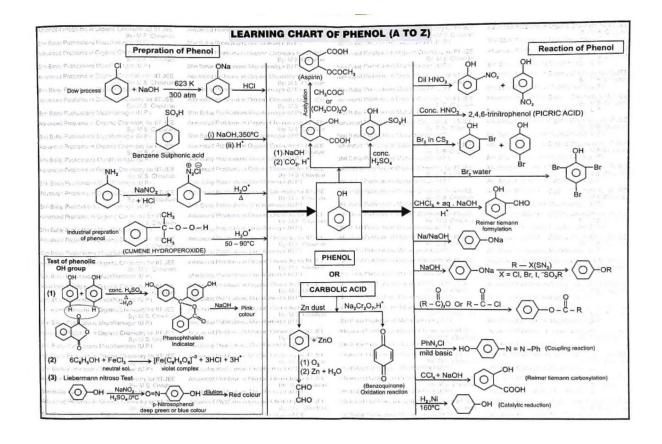
M.S. CHOUHAN

Contents

	1	General Organic Chemistry	1-72
	2	Isomerism (Structural & Stereoisomerism)	73-151
	3	Grignard Reagent	152-173
	4A	Hydrocarbons (Alkanes)	174-189
1	4B	Hydrocarbons (Alkenes)	190-253
8	4C	Hydrocarbons (Alkynes)	254-260
	5A	Alkyl Halides (Substitution Reaction)	261-297
1	5B	Alkyl Halides (Elimination Reaction)	298-328
	5C	Alkyl Halides	329-341
	6	Alcohol, Ether and Epoxides	342-388
	7	Aldehydes and Ketones	389-440
	8	Aldol and Cannizaro Reaction	441-460
9	9	Carboxylic Acid and their Derivatives	461-479
	10	Amines	480-486
	11	Carbene and Nitrene	487-506
	12	Aromatic Compounds	507-578
	13	Practical Organic Chemistry	579-584
	14	Biomolecules	585-596
	15	IUPAC Names	597-624
	•	Learning Chart	





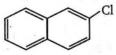


GENERAL ORGANIC CHEMISTRY



LEVEL- I

1. How many 2° Hydrogen atoms are present in the given following compound?



(a) 2

(b) 5

(c) 7

(d) 8

2. Identify which functional group is **Not** present in the given following compound?



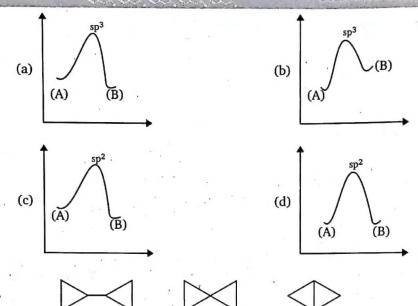
(a) Ketone

(b) Ester

(c) Amide

(d) Ether

3. Correct energy profile for amine inversion and hybridization of nitrogen in transition state is:



Correct order of the heats of combustion of above compounds is:

(a) (i)
$$>$$
 (ii) $>$ (iii) (

(i)

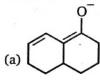
(ii)

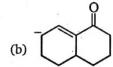
(c) (ii)
$$>$$
 (i) $>$ (iii)

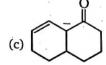
(iii)

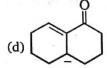
(d) (ii)
$$>$$
 (iii) $>$ (i)

Which of the following is not a resonance structure of the others?

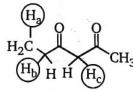








Rank the hydrogen atoms (Ha, Hb, Hc) present in the following molecule in decreasing order of their acidic strength.



(a)
$$a > b > c$$

(b)
$$b > a > c$$

(c)
$$b > c > a$$

(d)
$$c > b > a$$

The correct relation between the bond lengths a and b is:

(a)
$$a = b$$

(b)
$$b > a$$

(c)
$$b < a$$

8. The number of $sp^2 - sp^2$ sigma bonds in the compound given below is :



(a) 1

- (b) 3
- (c) 4
- (d) 5
- 9. The total number of lone pair of electrons in the given molecule is:



(a) 2

(b) 3

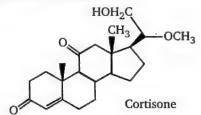
- (c) 4
- . (d) 5
- 10. Which of the following rings is highly strained?



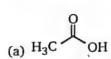
(b)

(c) β-lactone

The functional groups present in Cortisone are : 11.



- (a) ether, alkene, alcohol
- (b) alcohol, ketone, alkene, ether
- (c) alcohol, ketone, amine
- (d) ether, amine, ketone
- Select the acid with the highest K_a (i.e., lowest pK_a). 12.

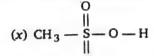


(d) C

Most stable carbocation among the following is: 13.



- (d) [⊕] CH₃
- Arrange the following in increasing order of their pK_a values. 14.



(y) CH₃ — C— O — H

(z) CH₃ — OH

- (b) x < y < z (c) y < z < x
- (d) x < z < y

15. Which is the major product of the following reaction?

$$(a) \qquad \begin{array}{c} H \\ N \\ N \\ H \end{array} \qquad \begin{array}{c} HCl \\ 1 \text{ mole} \end{array} \qquad Product$$

$$(b) \qquad \begin{array}{c} H \\ N \\ N \\ H \end{array} \qquad \begin{array}{c} H \\ Cl \\ H \end{array} \qquad \begin{array}{c} H \\ C$$

16. In the given pair identify most acidic compound in (A) and (B). Most basic in (C) and (D).

17. Several factors (steric, electronic, orbital interactions etc.) can affect the inversion barrier of an amine. In the given pair which data is correctly placed?

(a) Me

N

Me

N

Me

N

Me

N

Me

$$\Delta G^{\dagger} = 7.9 \text{ kcal/mol}$$

A

(b)
$$\Delta G^{\dagger} = 20.5 \text{ kcal/mol } \Delta G^{\dagger} = 7.0 \text{ kcal/mol}$$

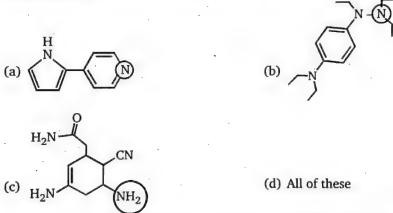
GENERAL ORGANIC CHEMISTRY

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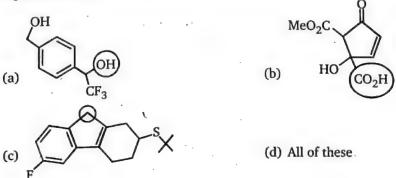
Me
$$V_{S}$$
 V_{S}
 V

- (d) All of these
- **18.** Select the response that correctly identifies the number of carbon atoms of each type of hybridization in the compound given below

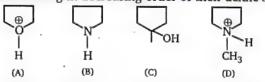
19. Circle represents most basic atoms in these molecule. Which of the following is correct representation?



20. Circle represent most acidic hydrogens in these molecules. Which of the following is correct representation?



Arrange the following in decreasing order of their acidic strengths. 21.



(a) A > C > B > D (b) A > D > B > C(c) A > D > C > B (d) D > A > C > B

22. Cyclopropane '

Cyclobutane Cyclopentane (II) (III)

The correct order of heats of combustion of above compounds is:

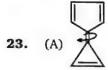
(a) I > II > III

(I)

III < I < II (d)

(c) III > II > I

II < I < III (b)







Compare carbon-carbon bond rotation across A, B, and C

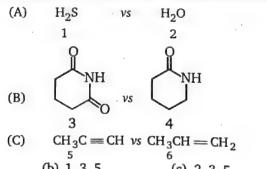
(a) A > B > C

(b) A > C > B

(c) B > A > C

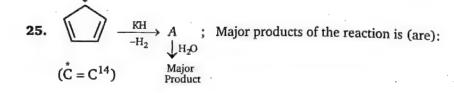
(d) B > C > A

Which of the following acids would have a STRONGER CONJUGATE BASE? 24.



(a) 2, 4, 6

- (b) 1, 3, 5
- (c) 2, 3, 5
- (d) 1, 3, 6



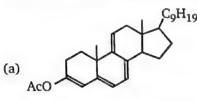


(b)

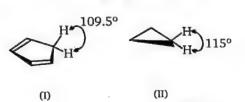
(c)

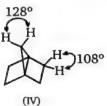
(d) both (b)& (c)

26. Which of the following compound is most stable?



27. Selected bond angles for six hydrocarbons are shown below. Arrange these hydrocarbons according to their pK_a values, from the lowest to the highest.





117° (III)

- (a) V < I < VI < II < III < IV
- (b) IV < I < II < III < V < VI
- (c) II < IV < I < VI < V < III
- (d) I < V < IV < III < II < VI
- 28. Which statement about the following equilibrium is true?

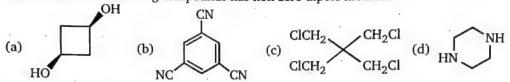
$$O^{-}K^{+} + H_{2}O \longrightarrow OH + K^{+}OH^{-}$$
t-butoxide $pK_{a} = 15.7$ $pK_{a} = 18$

- (a) The equilibrium favours the products
- (b) t-Butoxide is the dominant anionic species in the equilibrium
- (c) Water is the weaker acid
- (d) t-Butoxide is stabilized by resonance
- 29. Consider the following reaction involving two acids shown below: formic acid and HF.

$$K^{+}F^{-} + H \longrightarrow OH \longrightarrow H \longrightarrow O^{-}K^{+} + HF$$
 $pK_{a} = 3.8$

Which of the following statements about this reaction are true?

- (A) Formic acid is the strongest Bronsted acid in the reaction
- (B) HF is the strongest Bronsted acid in the reaction
- (C) KF is the strongest Bronsted base in the reaction
- (D) KO₂CH is the strongest Bronsted base in the reaction
- (E) The equilibrium favours the reactants
- (F) The equilibrium favours the products
- (G) Formic acid has a weaker conjugate base
- (H) HF has a weaker conjugate base
- (a) A, D and F
- (b) B, D, and H
- (c) A, C, and H
- (d) B, D, E and H
- 30. Which one of the following compounds has non zero dipole moment?



31. Which one of the following has the smallest heat of combustion?

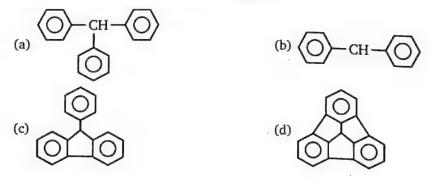
(a)
$$H_3C$$
 $C(CH_3)_3$ (b) H_3C $C(CH_3)_3$ (c) H_3C $C(CH_3)_3$ (d) H_3C $C(CH_3)_3$

32. Rank the following substances in order of decreasing heat of combustion (maximum \rightarrow minimum).

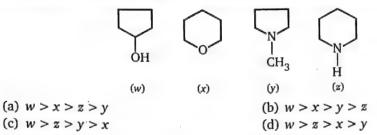
CH₃

$$1$$
(a) $1 > 2 > 4 > 3$
(b) $3 > 4 > 2 > 1$
(c) $2 > 4 > 1 > 3$
(d) $1 > 3 > 4 > 2 > 4$

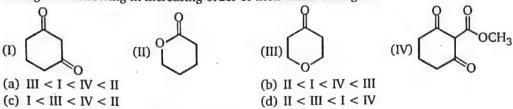
33. Which of the following has lowest pK_a value?



34. Arrange the following (w, x, y, z) in decreasing order of their boiling points:



35. Arrange the following in increasing order of their acidic strength.

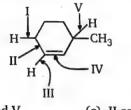


36. How many degrees of unsaturation are there the following compound?

37. The heat of hydrogenation for 3-methylbutene and 2-pentene are -30 kcal/mol and -28 kcal/mol respectively. The heats of combustion of 2-methylbutane and pentane are - 784 kcal/mol and -782 kcal/mol respectively. All the values are given under standard conditions. Taking into account that combustion of both alkanes give the same products, what is ΔH (in kcal/mol) for the following reaction under same conditions?

(a) 0 (b)
$$-4$$
 (c) -2 (d) 2

38. Which of the following σ -bonds participate in hyperconjugation ?



(a) I and II

(a) 6

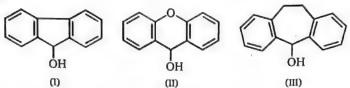
(b) I and V

(c) II and V

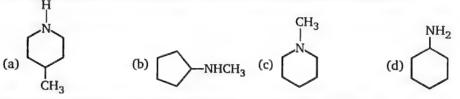
(d) III and IV

Decreasing order of acidic strength of different (-OH) groups is:

- (a) w > x > y > z
- (b) w > z > x > y
- (c) z > w > x > y
- (d) z > x > w > y
- 40. Arrange the following alcohols in decreasing order of the ease of ionization under acidic conditions.



- (a) I > III > II
- III < II < I
- (c) II > III > I
- (d) II > I > III
- Among the isomeric amines select the one with the lowest boiling point. 41.



- Which one of the compounds shown below, is not an isomer of the others?
 - (a) (b)
- (d)
- Arrange the anions (p) $\overline{C}H_3$, (q) $\overline{N}H_2$, (r) OH^- , (s) F^- , in decreasing order of their basic 43. strength.
 - (a) p > q > r > s
- (b) q > p > r > s
- (c) r > q > p > s
- (d) r > p > q > s
- 44. One among the following compounds will not give effervescence with sodium carbonate:
 - SO_3H NO-(a) C₆H₅CO₂H (c) C_6H_5OH (d)
- 45. The carboxylic acid which has maximum solubility in water is:
 - (a) phthalic acid

(b) succinic acid

(c) malonic acid

(d) salicylic acid

46. Among the following compounds, the most basic compound is :



(b) N

(c) (



47. ×

Arrange the (C - H) bonds x, y and z in decreasing order of their bond dissociation energies in homolysis.

- (a) y > x > z
- (b) z > x > y
- (c) z > y > x
- (d) y > z > x
- 48. 23 g of sodium will react with methyl alcohol to give:
 - (a) one mole of oxygen

(b) 22.4 dm³ of hydrogen gas at NTP

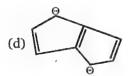
(c) 1 mole of H₂

- (d) 11.2 L of hydrogen gas at NTP
- **49.** Which of the following is most polar?









50. NH; N-H; N

The correct order of decreasing basic strengths of x, y and z is:

- (a) x > y > z
- (b) x > z > y
- (c) y > x > z
- (d) y > z > x
- 51. Which of the following is the strongest Bronsted acid?









52. Which of the following is the strongest Bronsted base?



(b) N





- 53. Which of the following is polar aprotic solvent?
 - (a) DMSO
- (b) Crown ether
- (c) DMG
- (d) All of these

- 54. Some pairs of acids are given below. Select the pair in which second acid is stronger than first
 - (a) CH₃CO₂H and CH₂FCO₂H
 - (b) CH2FCO2H and CH2ClCO2H
 - (c) CH2ClCO2H and CH2BrCO2H
 - (d) CH3CH2CHFCO2H and CH3CHFCH2CO2H
- **55.** $H C \equiv C C \equiv C CH_3$;

Compare the bond lengths a and b:

- (a) a = b
- (b) a > b
- (c) b > a
- (d) a >>> b
- 56. Which (isomeric) amine has lowest boiling point?
 - (a) 1º amine

(b) 2° amine

(c) 3° amine

- (d) cannot predict
- - (a) (2-)
 - (c)

- (b) (2+) 2SbCl₆
- (d) mixture of (a) and (b)
- 58. Which of the following substances is not an isomer of 3-ethyl 2-methyl pentane?
 - (a) (c)
- (b)
- (d) All are isomers
- 59. Which of the following is an isomer of compound 1?
 - CH₃CH₂CHO CH₃ CH₃ CH₃ CH = CH- OH CH₂ CH- CH₃ (1) (2) (3)
 - (a) 2
- (b) 4
- (c) 2 and 3
- (d) all are isomers

 $60. \qquad \stackrel{\text{Br}}{ } \xrightarrow{\text{AgNO}_3} (A)$

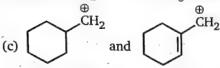
Which statement is incorrect in respect of the above reaction?

(a) Product is aromatic

- (b) Product has high dipole moment
- (c) Product has less resonance energy
- (d) Product is soluble in polar solvent

GENERAL ORGANIC CHEMISTRY

- 61. Some pairs of ions are given below. In which pair, first ion is more stable than second?
 - (a) $CH_3 \overset{\oplus}{C}H CH_3$ and $CH_3 \overset{\oplus}{C}H OCH_3$
 - (b) $CH_3 CH_2 \overset{\oplus}{C}H CH_3$ and $CH_2 = CH CH_2 \overset{\oplus}{C}H_2$



- $\begin{array}{c|ccccc} \operatorname{CH_3} \operatorname{CH} \operatorname{CH_3} & \operatorname{CH_3} \operatorname{N} & -\operatorname{CH_3} \\ \text{(d)} & & & & & & & \\ \operatorname{CH_2} \operatorname{C_{\oplus}} \operatorname{CH_3} & & & \operatorname{CH_3} \operatorname{C}^{\oplus} \operatorname{CH_3} \\ \end{array}$
- 62. Among the given pairs in which pair, first compound has higher boiling point than second?
 - (a) $CH_3 CH_2OCH_3$ and $CH_3 CH CH_3$

$$\begin{array}{c} \text{OH} \\ \text{(b) } \text{CH}_3 - \text{CH}_2 - \text{CH}_2 - \text{CH}_3 \text{ and } \text{CH}_3 - \text{CH}_2 - \text{CH}_3 \\ & \text{CH}_2 - \text{CH}_3 \\ \text{(c) } \text{CH}_3 - \text{CH}_2 - \text{CH}_2 - \text{CH}_2 - \text{CH}_3 \text{ and } \text{CH}_3 - \text{CH} - \text{CH}_2 - \text{OH} \\ \text{(d) } \text{CH}_3 - \text{CH}_2 - \text{CH}_2 - \text{CH}_3 \text{ and } \text{CH}_3 - \text{CH}_2 - \text{CH}_2 - \text{CH}_2 \end{array}$$

- 63. Which of the following alcohols is the least soluble in water?
 - (a) Ethanol

(b) 1-Propanol

(c) 1-Butanol

- (d) 1-Pentanol
- **64.** Which of the following alcohols is expected to have a lowest pK_a value?
 - (a) Ethanol

- (b) 1-propanol
- (c) 2, 2, 2-trifluorethanol
- (d) 2-chloroethanol
- 65. Which of the following alkenes is the most stable?

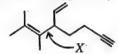


(b)





66. Bond X is made by the overlap of which type of hybridized orbitals?



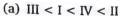
(a) sp and sp^3

(b) sp and sp^2

(c) sp^2 and sp^3

- (d) none of these
- 67. Increasing order of acidic strength of given compounds is:

$$H_3C$$
 (I) NC (III) H_3CO (III) (IV)



(c) I < III < IV < II

(b) II > I > IV < III

(d)
$$I < III < II < IV$$

68. \bigcirc COOH + NaHCO₃ \longrightarrow CO₂ + \bigcirc COONa, $\stackrel{\star}{C}$ is with the product :

(a) CO₂

(b) COONa (c) both

(d) none of these

69. Rank in the order of increasing acidity.



(II) NH



(a) III < I < II

(c) III < II < I

(b)
$$I < III < II$$

(d) II < I < III

70. Which compound has the highest value of pK_a ?

(a)
$$Cl - CH_2 - CH_2 - COOH$$

71. Consider the hydrogen atoms attached to three different carbon atoms (labeled 1, 2 & 3). Rank the attached hydrogen atoms in order from most acidic to least acidic.

$$\bigcap_{1}^{9} \bigcap_{2}^{3}$$

(a) 2 > 1 > 3

(b) 1 > 2 > 3

(c) 2 > 3 > 1

(d) 3 > 2 > 1

72. Decreasing order of acidic strengths of following compounds is:



(a) x > y > 2



(b) y > x > z

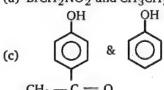


(c) z > y > x

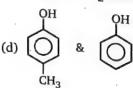
(d) z > x > y

73. Among the given pairs, in which pair second compound is more acidic than first?

(a) BrCH₂NO₂ and CH₃CH₃

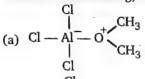


(b) $CH_3 - CCH_2CN$ and $CH_3 - C-CH_3$



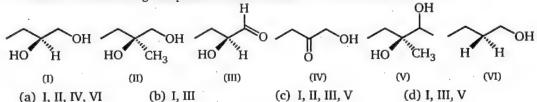
- Which of the underlined atoms in the molecules shown below have sp-hybridization?
 - (u) CH2CHCH3
- (v) CH2CCHCl
- (w) $CH_3CH_2^+$
- (x) $H C \equiv C H$

- (y) CH₃CN (a) x and z
- (z) $(CH_3)_2CNNH_2$ (b) x, y, and z
 - (c) u, w and x
- (d) v, x and y
- Which of the following, is the product of the reaction between $AlCl_3$ and CH_3OCH_3 ? 75.

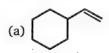


(c)
$$Cl - Al^{+} - O$$
 CH_{3} CH_{3}

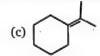
Which of the following compounds contain at least one secondary alcohol?



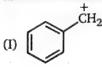
- (c) I, II, III, V
- (d) I, III, V
- Which of the following has the most negative heat of hydrogenation?





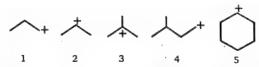


- (d)
- Which of the following options is the correct order of relative stabilities of cations I, II and III 78. as written below (most stable first)?



- (II) $H_2C = CH CH_2 CH CH_3$
- CH₃

- (a) I > II > III
- (b) II > III > I
- (c) III > I > II
- (d) I > III > II
- What is the decreasing order of stability (most stable → least stable) of the following 79. carbocations?



(a) 3 > 2 > 1 > 4 > 5

(b) 3 > 2 > 5 > 4 > 1

(c) $1 \approx 4 > 2 \approx 5 > 3$

(d) $3 > 1 \approx 4 > 2 \approx 5$

80.

16

the hydrogen indicated by arrow will be easily removed as:

- (b) H^Θ
- (c) H*
- (d) H^{-2}
- Rank the bond dissociation energies of the bonds indicated with the arrows. (from smallest 81. to largest).

$$\begin{array}{c} \stackrel{1}{\underset{H}{\bigvee}} \stackrel{H}{\underset{CH_2}{\bigvee}} 2$$

- (a) 1 < 2 < 3
- (b) 3 < 2 < 1
- (c) 2 < 3 < 1
- (d) 3 < 1 < 2
- 82. Rank the following compounds in order of decreasing acid strength (most acidic → least acidic).

$$\bigcirc_{NO_2}^{OH} \bigcirc_{NO_2}^{OH} \bigcirc_{NO_2}^{OH}$$

- (a) 2 > 4 > 1 > 3 (b) 1 > 3 > 4 > 2
- (c) 3 > 1 > 2 > 4
- (d) 3 > 1 > 4 > 2
- Rank the following compounds in order of increasing acidity (weakest acid first). 83.

- (a) 2 < 3 < 1
- (b) 3 < 1 < 2
- (c) 1 < 2 < 3
- (d) 2 < 1 < 3
- Which of the following phenols has the largest pK_a value (i. e., is least acidic)? 84.

(b) O₂N -

(c) H₃C —

- (d) N≡C -
- Among the given sets, which represents the resonating structures? 85.

(a)
$$H - C \equiv N - 0$$
: and $H - 0 - C \equiv N$:

(b)
$$H - \overset{+}{0} = C = \overset{-}{N}$$
: and $H - \overset{-}{0} - C = N$:

(c)
$$H - C = \stackrel{+}{N} - \stackrel{\cdots}{0}$$
; and $H - C - \stackrel{\cdots}{N}$; (d) $H - \stackrel{\cdots}{0} - C = \stackrel{\cdots}{N}$; and $H - \stackrel{\cdots}{N} = C = \stackrel{\cdots}{0}$;

(d)
$$H - \ddot{O} - C = N$$
 and $H - \ddot{N} = C = \ddot{O}$:

86. Identify each species in the following equilibrium according to the code:

SA = stronger acid; SB = stronger base; WA = weaker acid; WB = weaker base.

The pK_a of $(CH_3)_2NH$ is 36; the pK_a of CH_3OH is 15.2.

$$CH_3OH + (CH_3)_2NH \rightleftharpoons CH_3 - O^- + CH_3 - NH - CH_3$$

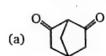
- 1 2
- 1 2
- 1 2
- 1 2

- (a) WA WB
- (b) WB WA
- (c) SA SB
- (d) SB SA

- (e) WA WA
- 87. The hydrogen bonding is strongest in which one of the following set?
 - (a) F --- F
- (b) O H - S
- (c) S H - F
- (d) F -- H --- O
- 88. Intermolecular hydrogen bonding is strongest in :
 - (a) methylamine
- (b) phenol
- (c) formaldehyde
- (d) methanol

Identify most acidic hydrogen in given compound.

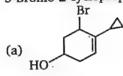
- (a) a
- (b) b
- (c) c
- (d) d
- 90. Which of the following compounds would you expect to be strongest carbon acid?

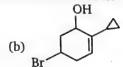




(c) $CH_2(CO_2Et)_2$

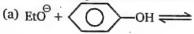
- (d) CH₃COCH₂COOC₂H₅
- 91. 5-Bromo-2-cyclopropyl cyclohex-2-enol have correct structure is:



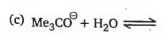


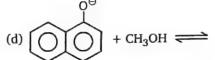
- **92.** Rearrange the following in the increasing order of acidic strength.
 - (i) benzoic acid
- (ii) p-methoxybenzoic acid
- (iii) o-methyoxybenzoic acid

- (a) i < ii < iii
- (b) iii < i < ii
- (c) ii < i < iii
- (d) iii < ii < i
- 93. In the following acid-base reaction, in which can backward reaction if favoured?



(b) KH + EtOH





- 94. Which compound posses highest dipole moment?
 - (a) naphthalene

(b) phenanthrene

(c) anthracene

(d) azulene

(E = activation energy)

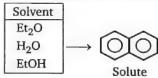
Relation between activation energies of above reactions is:

(a)
$$E_2 > E_1 > E_3$$

(b)
$$E_3 > E_1 > E_2$$

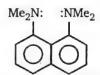
(c)
$$E_3 > E_2 > E_1$$

- (d) $E_1 > E_2 > E_3$
- 96. Rank the following solvents in decreasing order of ability to dissolve given compound.



- (a) $Et_2O > H_2O > EtOH$
- (b) $H_2O > EtOH > Et_2O$
- (c) $H_2O > Et_2O > EtOH$

(d) $Et_2O > EtOH > H_2O$



97.

1, 8-Bis (dimethylamino) naphthalene is after referred so as (Proton sponge)

Its basic strength is 1010 more than 1-dimethyl amino naphthalene.

Reason for high basic strength is:

(a) resonance

(b) steric inhibitation of resonance

(c) ortho effect

(d) hyperconjugation

98. In the given pair of compounds, in which pair second compound has higher boiling point than first compound?

(a) OH and O

(b) O and

(c) $HO - CH_2 - CH_2 - OH$ and $CH_3 - CH_2 - CH_2 - OH$

(d) $\stackrel{\text{CH}_3}{\stackrel{\mid}{\bigcap}}$ and $\stackrel{\text{NH}}{\stackrel{\mid}{\bigcap}}$

99. Me Me Me Me Me Me Me Me

Me

Me

Dipole moments of given compound will be:

- (a) (A) = 6.87D, (B) = 4.11D
- (b) (A) = 4.11 D, (B) = 6.87 D
- (c) (A) = 4.11 D, (B) = 4.11 D
- (d) (A) = 6.87 D, (B) = 6.87 D
- 100. Order of decreasing basic strengths of halides is:
 - (a) $F^- > Cl^- > I^- > Br^-$

(b) $F^- > Cl^- > Br^- > I^-$

(c) $I^- > Br^- > Cl^- > F^-$

- (d) $I^- > Cl^- > Br^- > F^-$
- 101. Among the xylenes, which is thermodynamically most stable?

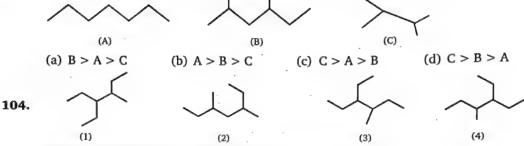
(a) (1)

(p) (C)



- (d) All are equally stable
- **102.** Heat of combustion of two isomer *x* and *y* are 17 kJ/mol and 12 kJ/mol respectively. From this information it may be concluded that :
 - (a) isomer x is 5 kJ/mol more stable
 - (b) isomer y is 5 kJ/mol less stable
 - (c) isomer y has 5 kJ/mol more potential energy
 - (d) isomer x is 5 kJ/mol less stable

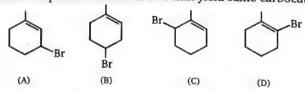
Rank the following substances in decreasing order of heat of combustion (most exothermic 103. least exothermic)



Choose the statement that best describes given compounds.

- (a) 1, 3, 4 represent same compound
- (b) 1 and 3 are isomer of 2 and 4
- (c) 1, 4 are isomer of 2 and 3
- (d) All the structure represent the same compound
- 105. Decreasing order of acid strengths is:

- Rank the following in decreasing order of basic strength is: 106.
 - (A) $CH_3 CH_2 C \equiv C$
- (B) $CH_3 CH_2 S^-$
- (C) $CH_3 CH_2 CO_2^-$
- (D) $CH_3 CH_2 O^-$
- (a) B > A > D > C (b) D > A > B > C
- (c) A > D > B > C (d) A > D > C > B
- Among the given compound choose the two that yield same carbocation on ionization. 107.



- (a) A, C
- (b) B, D
- (c) A, B
- (d) B, C

108. Oxalic acid pK_1

Malonic acid

 pK_2

Heptanedioic acid pK_3

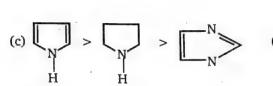
where pK_1 , pK_2 , pK_3 are first ionization constants. Incorrect order is:

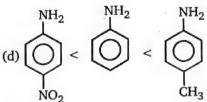
(a) $pK_1 > pK_2 > pK_3$ (b) $pK_1 < pK_2 < pK_3$ (c) $pK_3 > pK_2 > pK_1$ (d) $pK_3 > pK_1 > pK_2$

In sets a – d, only one of the set is incorrect regarding basic strength. Select it: NH_2

(a)
$$Ph - NH - Ph_1 < Ph - NH_2 <$$
 (strong base)

$$(b) \begin{array}{|c|c|} \hline \\ N \\ \hline \end{array} < \begin{array}{|c|c|} \hline \\ N \\ \hline \end{array} < \begin{array}{|c|c|} \hline \\ N \\ \hline \end{array}$$





110. Dipole moment of which ketone is maximum?









- 111. Correct order of basic strengths of given amines is:
 - (a) $Me_2NH > MeNH_2 > Me_3N > NH_3$ (Protic solvent)
 - (b) $\text{Et}_{2}^{NH} > \text{Et}_{3}^{H} > \text{Et}_{1}^{NH}_{2} > \text{NH}_{3}$ (Protic solvent)
 - (c) $Me_3N > Me_2NH > Me-NH_2 > NH_3$ (Gas phase)
 - (d) All are correct
- 112. Order of basic strength Ph NH $_2$, Ph NH- Me, Ph N- Me, Me

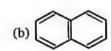
 (A)

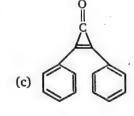
 (B)

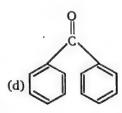
 (C)

 (D)
 - (a) A > B > C > D (b) B > A > C > D (c) C > B > A > D (d) C > B > D > A
- 113. Carbon-carbon double bond length will be maximum in which of the following compounds?
 - (a) $CH_3 CH = CH_2$ (c) $CH_3 - C = C - CH_3$ | | $CH_3 CH_3$
- (b) $CH_3 CH = CH CH_3$
- (d) $CH_2 = CH_2$
- 114. Which has maximum dipole moment?

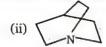








115. (i) Et₃N



(iii)

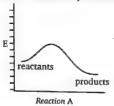
Compare the basic strengths of compounds given:

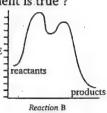
(a) (i)
$$>$$
 (ii) $>$ (iii)

(a) (i)
$$>$$
 (ii) $>$ (iii) (b) (ii) $>$ (i) $>$ (iii) (c) (ii) $>$ (ii) $>$ (i) (d) (iii) $>$ (i) $>$ (i)

(d) (iii)
$$>$$
 (ii) $>$ (i

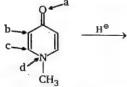
116. For the following two reactions, which statement is true?





- (a) Reaction A is faster and less exergonic than B
- (b) Reaction B is faster and more exergonic than A
- (c) Reaction A is faster and less endergonic than B
- (d) Reaction B is faster and more endergonic than A

117.



Identify the site, where attack of H+ is most favourable.

Rank the following alkenes on order of increasing λ_{max} 118.







(a)
$$1 < 2 < 3$$

(b)
$$1 < 3 < 2$$

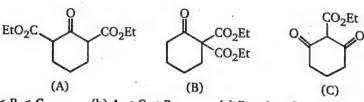
Which of the following cyclic amine has lowest ΔG^* for inversion?

(a)
$$N - CH_3$$
-Me (b) $N - n$ -

(c)
$$N - t$$
-Bu

$$(d)$$
 $N - Ph$

120. Rank in the order of increasing acidic strength:



(a) A < B < C

(b) A < C < B

(c) B < A < C

(d) B < C < A

Which one of the following dienes would you expect to be the most stable?

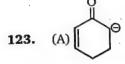








- 122. Which metal catalyzed reaction would release the maximum amount of heat per CH_2 unit?
 - (a) cyclopropane $+H_2 \rightarrow propane$
- (b) cyclobutane $+H_2 \rightarrow$ butane
- (c) cyclopentane $+H_2 \rightarrow pentane$
- (d) cyclohexane $+H_2 \rightarrow$ hexane







Compare basic strengths of the above compounds:

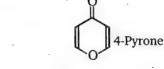
(a)
$$A > B > C$$

(b)
$$B > A > C$$

(c)
$$C > A > B$$

(d)
$$C > B > A$$

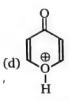
124. On reaction with acid, 4-pyrone gives a very stable cationic product. Which of the following structures shows the protonation site in that product?



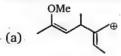


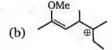


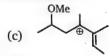


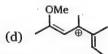


125. Which of the following is the most stabilized carbocation?

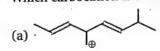


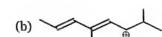


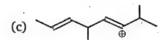




126. Which carbocation is the most stable?







- **127.** Consider a positively charged C_2H_3 species in which the positively charged carbon is sp-hybridized, the uncharged carbon is sp^2 -hybridized and an empty p-orbital is perpendicular to the π system. What it the best description of this cation ?
 - (a) vinyl
- (b) allenyl
- (c) alkyl
- (d) allyl

128. Which of the following reactions is not exothermic?

(a)
$$CH_3 - CI + CH_3 - CH_3 \longrightarrow CH_4 + CH_3 - CH_2 - CI$$

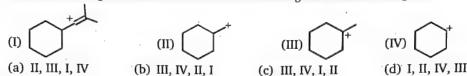
(b)
$$CH_3 - CI + (CH_3)_3 C - H \longrightarrow CH_4 + (CH_3)_3 C - CI$$

(b)
$$CH_3 - CI + (CH_3)_3 C - H \longrightarrow CH_4 + (CH_3)_3 C - CI$$

(c) $CH_3 - CI + CH_2 = CH - CH_3 \longrightarrow CH_4 + CH_2 = CH - CH_2 - CI$
(d) $CH_3 - CI + CH_3 - CH_3 - CH_4 + CH_2 = CH_3 - CH_3 - CH_4 + CH_2 - CH_3 - CH_4 + CH_3 - CH_4 - CH_3 - CH_4 - CH_5 - CH_5$

(d)
$$CH_3 - Cl + CH_2 = CH_2$$
 \longrightarrow $CH_4 + CH_2 = CHCl$

129. List the following carbocations in order of decreasing stabilization energies.



130. For the following two acid-base reactions, which statement is true?

(I)
$$CH_3CH_2^- + CH_3NH_2 \atop pK_a = 35$$
 \Longrightarrow $CH_3CH_3 + CH_3NH_2 \atop pK_a = 50$ (II) $F^- + H_2O \atop pK_a = 15.7$ \Longrightarrow $HF \atop pK_a = 3.2$ $+ HO^-$

- (a) I is favoured to the right, II is favoured to the left
- (b) I is favoured to the left, II is favoured to the right
- (c) I is favoured to the right, II is favoured to the right
- (d) I is favoured to the left, II is favoured to the left

Rank the hydrogen atoms (H_a, H_b, H_c) in the following molecules according to their acidic 131. strengths:

(a)
$$a > b > c$$

(b)
$$b > a > c$$

(c)
$$b > c > a$$

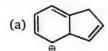
(d)
$$a > c > 1$$

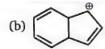
In which of the following reactions, backward reaction is favoured?

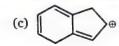
(a)
$$H - C \equiv C - H + Li^{+} - CH_{2}CH_{3}$$
 \rightleftharpoons $H - C \equiv C : {}^{\Theta}Li^{+} + H_{3}C - CH_{3}$
(b) $F_{3}C$ OH $+$ $^{-}OCH_{2}CH_{3}$ \rightleftharpoons $F_{3}C$ $O\overline{:}$ $+$ $^{+}HOCH_{2}CH_{3}$
(c) $CH_{3}CH_{2}\overset{+}{S}H_{2} + CH_{3}CH_{2}OH$ \rightleftharpoons $CH_{3}CH_{2}SH + CH_{3}CH_{2}\overset{\oplus}{O} - H$

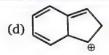
133. Which carbocation is the most stabilized?

GENERAL ORGANIC CHEMISTRY









134. Taking into account of hybridization and resonance effects, rank the following bonds in order of decreasing bond length.



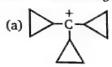
(a) I > II = III

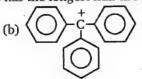
(b) II > III > I

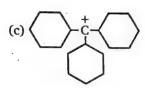
(c) I > III > II

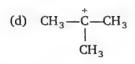
(d) $\Pi = \Pi = I$

135. Which one among the following carbocations has the longest half-life?

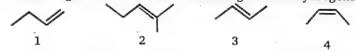








Rank the following alkenes in order of decreasing heats of hydrogenation (largest first) 136.

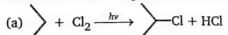


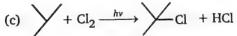
(a) 2 > 3 > 4 > 1

(b) 2 > 4 > 3 > 1 (c) 1 > 3 > 4 > 2

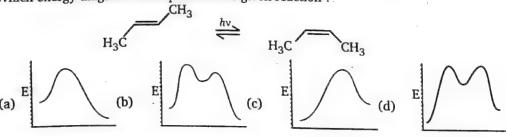
(d) 1 > 4 > 3 > 2

Which of the following reactions is most exothermic? 137.



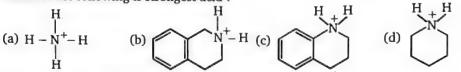


Which energy diagram best represents the given reaction? 138.

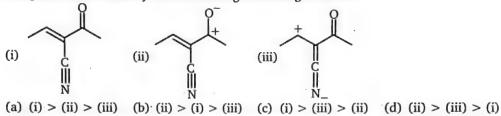


139. Which one of the following is most stable?

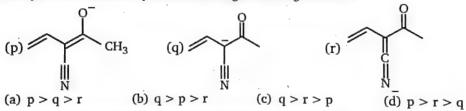
140. Which of the following is strongest acid?



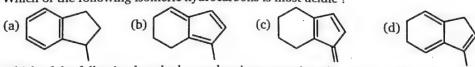
141. Compare relative stability of the following resonating structure.



142. Compare relative stability of the following resonating structure.



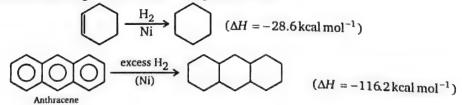
143. Which of the following isomeric hydrocarbons is most acidic?



144. Which of the following has the lowest barrier to rotation about the indicated bond?



145. Use the following data to answer the question below.



(a) 84 kcal/mol

- (b) 100 kcal/mol
- (c) 110 kcal/mol
- (d) 116 kcal/mol .
- **146.** How many double bond equivalents does a compound of molecular formula C₆H₁₂O₆ possess?

(a) 0

- (b) 1
- (c) 2
- (d) 3
- 147. How many double bond equivalents does amoxycillin (shown below) possess?

(a) 5

(b) 6

(c) 7

(d) 9

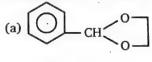
148. What is the oxidation state of osmium in 7B and 7C, respectively?

(a) 6, 8

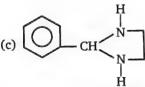
149.

Identify most acidic hydrogen present in the above compound:

- (a)
- (b) h
- (c) c
- (d) d
- 150. Which of the following compounds has most acidic hydrogen?



(b) CH S

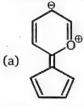


- **151.** Acetic acid, (CH₃COOH), has a p K_a of 4.8. Ethanol, (CH₃CH₂OH), has a p K_a of 16.0. What are the major species present, when acetic acid and ethanol are added to water and the pH is adjusted to 7.0?
 - (a) CH₃CO₂H and CH₃CO₂OH
- (b) CH₃CH₂O⁻ and CH₃CH₂OH
- (c) CH₃CO₂H and CH₃CH₂O⁻
- (d) CH₃CO₂ and CH₃CH₂OH

152.

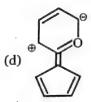


The most stable canonical structure of given molecule is:

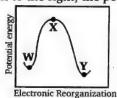








153. In the potential energy diagram to the right, the point X represents:



- (a) a transition state
- (c) a resonance hybrid

(b) a reaction intermediate

(d) a reactant

 $-CH_3$ 154. (I)





Which of the following orders is correct for heat of hydrogenation of these compounds?

- (a) I > III > II
- (b) III > II > I
- (c) III > I > II
- (d) II > I > III

 CH_3 155.





Which of the following orders is correct for heat of hydrogenation of these compounds?

- (a) I > II > III
- (b) III > II > I
- (c) II > III > I
- (d) III > I > II

 $CH_2 = O \longleftrightarrow {}^{\oplus}CH_2 - O^{\Theta} \longleftrightarrow {}^{\Theta}CH_2 - O^{\oplus}$ 156.

Which of these structures is practically not a valid canonical structure for formaldehyde?

- (b) II
- (c) III
- (d) None of these

 $CH_2 = CH - CH = CH - {}^{\oplus}NH_3$; 157.

$${}^{\oplus}$$
CH₂—CH =CH — ${}^{\ominus}$ CH — ${}^{\oplus}$ NH₃

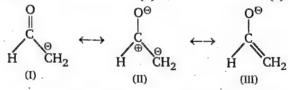
(I)
$$^{\oplus}$$
CH $_2$ - CH = CH - CH = NH $_3$ (III)

Which of these structures is not a valid canonical structure?

(a) I

- (b) II
- (c) III
- (d) none of these

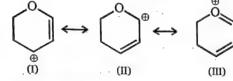
158.



The correct order of stability for the given canonical structures is:

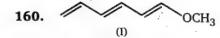
- (a) I > III > II
- II < I < III (d)
- (c) II > III > I
- (d) II > I > III

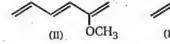




The most stable canonical structure among the given structure is:

- (b) II
- (c) III
- (d) all are equally stable

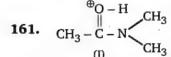


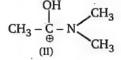




For the given compounds the correct order of resonance energy is:

- (a) III > I > II
- (b) II > I > III
- (c) I > II > III
- (d) III > II > I





$$CH_3 - C = N CH_3$$

$$CH_3 - CH_3$$

The correct stability order of the given canonical structures is :

- (a) I > II > III
- (b) III > I > II
- (c) I > III > II
- (d) II > III > I



In the above compound, how many sites are available for the attack of CH₂O⁻?

163. $CH_2 + CH_2$

- CH₃O CH + CH₂ CH₃O CH + CH C OEt

Which of the following orders of rotation barrier about the C = C bond, as indicated, is correct?

- (a) I > II > III
- (b) III > II > I (c) III > I > II
- (d) II > I > III

164.
$$\bigcirc$$
 CH \bigcirc CH \bigcirc CH \bigcirc CH \bigcirc CH \bigcirc OCH₃

Which of the following orders of rotation barrier about the C = C bond, as indicated, is correct?

- (a) I > II > III
- (b) II > II > I
- (c) III > I > II
- (d) II > I > III
- **165.** Which of the following compound is not resonance stabilized?



- (b) 🕞
- (c)
- (d) O

- **166.** Homologous compound have same:
 - (a) General formula
 - (c) Structural formula

- (b) Emperical formula
- (d) Molecular formula

167. Most acidic is:









- **168.** Which of the following substituents will decrease the acidic strength of phenol?
 - (a) -NO₂
- (b) -CN
- (c) —CH₃
- (d) -CHO
- 169. Which of the following structures possesses a cross-conjugated system?

(a)
$$CH_2 = CH - CH = CH - CH_2$$

(b)
$$CH_2 = CH - C = CH_2$$

(c)
$$CH_2 = CH - CH - CH = CH_2$$

 $CH = CH_2$

$$\begin{array}{c} {\rm CH_2CH_3} \\ {\rm (d)\ CH_2 = CH - C = CH_2} \\ {\rm | } \\ {\rm CH = CH_2} \end{array}$$

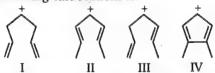
170. Examine the following resonating structures of formic acid for their individual stability and then answer the question given below.

$$\begin{array}{c} I \\ H - C - OH \longleftarrow H - C = O - H \longleftarrow H - C - OH \longleftarrow H - C - OH \\ III \\ III \\ III \\ III \\ III \\ IV \\ O^+ \\ O$$

Which of the following arrangements gives the correct order of decreasing stability of the above-mentioned resonance contributors?

- (a) II > I > III > IV
- (b) I > II > III > IV
- (c) IV > III > I > II
- II < I < III > I > II

- Which of the following is not resonating structure of each other?
 - (a) $CH_3 N = C = S$ and $CH_3 S C \equiv N$
 - (b) $CH_3 C = 0$ and $CH_3 C \equiv 0$
 - (c) $CH_3 C OH$ and $CH_3 C = O H$
 - (d) $CH_2 = CH C \equiv N$ and $CH_2 CH = C = N^-$
- In the molecule $CH_3C \equiv CCH = CH_2$, the maximum number of carbon atoms arranged 172. linearly is:
 - (a) 2
- (b) 3
- (c) 4
- (d) 5
- 173. The stability order of the following carbocations is:



- (a) II > IV > III > I (b) IV > II > III > I
- (c) II > III > I > IV
- (d) I > III > II > IV
- 174. Total number of α-hydrogen in given compound is:



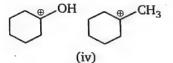
- (a) 4
- (b) 5
- (c) 6
- (d) 7
- In which pair second ion is more stable than first? 175.



(ii)



(iii)



- (a) (i) and (ii)
- (b) (ii) and (iii)
- (c) (ii) and (iv)
- (d) (iii) and (iv)
- Which one is the most stable cation in the following? 176.







The most reactive amine towards dilute hydrochloric acid is 177. CONH₂



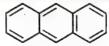
(b)





32

178. How many resonance structures are there for anthracene?



(a) 6

(b) 5

(c) 4

(d) 2

179. Which base is strong enough to convert (CH₃)₃COH into (CH₃)₃CONa in a reaction that goes to completion?

(a) NaNH₂

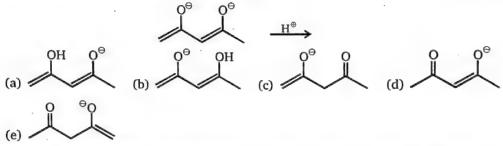
(b) CH₃CH₂Na

(c) NaOH

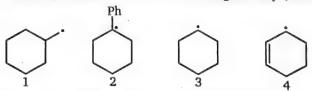
(d) CH₃CO₂Na

(e) More than one of the above

180. Based upon an understanding of product stability, predict the product formed when the following dianion reacts with one equivalent of acid



181. Rank the following alkyl radicals in order of increasing stability (least < < < most).



(a) 4 < 2 < 1 < 3

(b) 3 < 1 < 2 < 4

(c) 1 < 3 < 4 < 2

(d) 2 < 4 < 3 < 1

182. Among the given cations, the most stable carbonium ion is?

(a) sec-butyl

(b) tert-butyl

(c) n-butyl

(d) None of these

183. Cyclohexadiene contains _____ vinylic and _____ allylic hydrogen atoms ?



(a) 2 and 2 respectively

(b) 4 and 4 respectively

(c) 2 and 4 respectively

(d) 4 and 2 respectively

- 184. The dipole moments of halo compounds are in the order
 - (a) $CHCl_3 > CCl_4 > CHCl_2 > cis CHCl = CHCl$
 - (b) $cis CHCl = CHCl > CHCl_3 > CH_2Cl > CCl_4$
 - (c) cis CHCl = CHCl > CH₂Cl₂ > CHCl₃ > CCl
 - (d) CHCl₃ > CHCl₂ > cis CHCl = CHCl > CCl₄
- 185. The pka value in H₂O of picric acid, acetic acid and phenol are in the order:
 - (a) Picric acid 0.4, acetic acid 4.75, phenol 10.0

- (b) Acetic acid 0.4, picric acid 4.75, phenol 10.0
- (c) Picric acid 0.4 phenol 4.75, acetic acid 10.0
- (d) Phenol 0.4, acetic acid 4.75 picric acid 10.0
- **186.** The preferred sites of protonation in the following compounds are:

- (a) 1 and 3
- (b) 2 and 4
- (c) 1 and 4
- (d) 2 and 3

187. Among i-iii

$$(i) \qquad (ii) \qquad (iii) \qquad (iii) \qquad (iii)$$

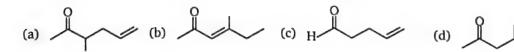
the boiling point follows the order

- (a) (ii) < (i) < (iii) (b) (iii) < (ii) < (i) (c) (i) < (ii) < (iii) (d) (ii) < (iii) < (i)
- 188. The number of C C sigma bonds in the compound

- (a) 16
- (b) 14
- (b) 18
- (d) 11
- 189. The correct order of dipole moment for the following molecules is

- (a) IV > I > III > II (b) I > IV > III > II (c) III > I > II > IV (d) II > III > IV > I
- **190.** Curved arrows are used in Organic Chemistry to show the movement of electrons in the mechanism of a reaction. The correct product of the following reaction is





~						ANSW	ERS	— LE	VEL 1	je					
1.	(c)	2.	(d)	3.	(d)	4.	(a)	5.	(d)	6.	(d)	7.	(b)	8.	(c)
9.	(b)	10.	(c)	11.	(b)	12.	(b)	13.	(a)	14.	(b)	15.	(b)	16.	(b)
17.	(d)	18.	(c)	19.	(d)	20.	(d)	21.	(c)	22.	(c)	23.	(c)	24.	(a)
25.	(d)	26.	(d)	27.	(d)	28.	(a)	29.	(d)	30.	(a)	31.	(c)	32.	(c)
33.	(d)	34.	(d)	35.	(d)	36.	(d)	37.	(b)	38.	(b)	39.	(a)	40.	(c)
41.	(c)	42.	(d)	43.	(a)	44:	(c)	45.	(c)	46.	(d)	47.	(b)	48.	(d)
49.	(b)	50.	(b)	51.	(d)	52.	(d)	53.	(d)	54.	(a)	55.	(c)	56.	(c)
57.	(b)	58.	(b)	59.	(d)	60.	(c)	61.	(b)	62.	(b)	63.	(d)	64.	(c)
65.	(d)	66.	(c)	67.	(a)	68.	(a)	69.	(d)	70.	(b)	71.	(a)	72.	(d)
73.	(d)	74.	(d)	75.	(a)	76.	(d)	77.	(a)	78.	(a)	79.	(b)	80.	(a)
81.	(d)	82.	(d)	83.	(d)	84.	(c)	85.	(b)	86.	(a)	87.	(a)	88.	(b)
89.	(a)	90.	(d)	91.	(b)	92.	(c)	93.	(d)	94.	(d)	95.	(d)	96.	(d)
97.	(b)	98.	(d)	99.	(a)	100.	(b)	101.	(b)	102.	(d)	103.	(a)	104.	(a)
105.	(c)	106.	(c)	107.	(c)	108.	(b)	109.	(c)	110.	(c)	111.	(d)	112.	(c)
113.	(c)	114.	(c)	115.	(c)	116.	(a)	117.	(a)	118.	(d)	119.	(c)	120.	(c)
121.	(c)	122.	(a)	123.	(c)	124.	(c)	125.	(d)	126.	(b)	127.	(a)	128.	(d)
129.	(b)	130.	(a)	131.	(c)	132.	(d)	133.	(c)	134.	(d)	135.	(a)	136.	(d)
137.	(c)	138.	(d)	139.	(c)	140.	(c)	141.	(a)	142.	(d)	143.	(b)	144.	(c)
145.	(a)	146.	(b)	147.	(d)	148.	(b)	149.	(a)	150.	(b)	151.	(d)	152.	(b)
153.	(a)	154.	(a)	155.	· (c)	156.	(c)	157.	(c)	158.	(b)	159.	(c)	160.	(c)
161.	(b)	162.	(c)	163.	(a)	164.	(a)	165.	(c)	166.	(a)	167.	(d)	168.	(c)
169.	(d)	170.	(b)	171.	(a)	172.	(c)	173.	(c)	174.	(c)	175.	(b)	176.	(b)
177.	(c)	178.	(c)	179.	(e)	180.	(d)	181.	(c)	182.	(b)	183.	(b)	184.	(c)
185.	(a)	186.	(a)	187.	(a)	188.	(b)	189.	(b)	190.	(c)				



LEVEL-2

1. Rank in order of radical stability (1 = most stable).

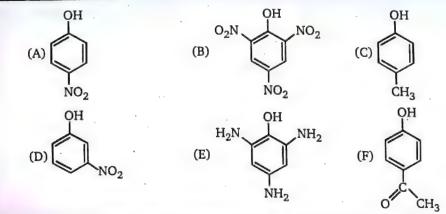
(a) (b) (c) (d) (d)

2. Predict the acidity order for the three phenols shown below : Acidity order : 1 (most) to 3 (least)

(a) Me OH (c) Me OH

Acidity order:

3. Comprehension



36			ORGANIC Chemistry for IIT-
A.	Which of the phenol deri	vatives above is the str	* ** * * *
	☐ Compound A☐ Compound D	☐ Compound B☐ Compound E	☐ Compound C☐ Compound F
В.	Which of the phenol deri	•	akest acid ?
	□ Compound A□ Compound D	☐ Compound B . ☐ Compound E	☐ Compound C☐ Compound F
C.	Which of the mono-nitro	•	e is the strongest acid?
	☐ Compound A	☐ Compound D	
D.	Which of the carbon-subs	tituted phenol derivati	ves above is the strongest acid?
	☐ Compound C	☐ Compound F	
4.	The following questions ras six choices in each ans	efer to the twelve compo	ounds given below. You may enter as a
	as six choices in each ans	wer box.	0
(a) N	(ь)	OH (c) $H_3C \longrightarrow O - C_2H$
(4	H—F	(e) CH	-CH ₃ (f)
G		(h)	о _{— Н} (i)
G	NH ₂	(k) H ₃ C —	
Α.	Which compound may ser		
В.	Which may serve both as F		
C.	Which compounds will no		
5.	Consider the following co	mpounds and answer A	and B.
	(I) (II)	N (III)	
	Which of the compounds i	s the strongest Property	Ĥ
A.	which of the commitment	S THE SHOHVEST Bronste	d acid ?

B. Which of the compounds is the strongest Lewis base?

(a) 1

(b) II

(c) III

(d) IV

6. Rank the non-bonding electrons indicated by the arrows in order of increasing energy.

7. In each of the following sections four nitrogen containing compounds are listed. In the box under each formula write a number (1 to 4) indicating the order of base strength.

un	der each formula write	e a number (1 to 4) indicating the order of	base strength.
(a)		$N-CH_3$	N-H	O ₂ N —NH ₂
(b)	н н	NH ₂	O ₂ N —NH ₂	N-H
(c)	N(CH ₃) ₂	N-H	N-H 	$N-CH_3$
(d)	H_3C $C \equiv N$		N-H	N-H

8. For the two sets of acids shown below, rank their acidity most acidic to least acidic.

$$(a) \xrightarrow{Br} OH \longrightarrow OH \longrightarrow OH \longrightarrow OH \longrightarrow OH$$

$$(ii) \longrightarrow OH \longrightarrow OH \longrightarrow OH$$

$$(iii) \longrightarrow OH \longrightarrow OH$$

$$(iii) \longrightarrow OH \longrightarrow OH$$

$$(iii) \longrightarrow OH$$

$$(iiii) \longrightarrow OH$$

$$(iiii) \longrightarrow OH$$

$$(iiii) \longrightarrow OH$$

$$(iv) \longrightarrow OH$$

9. In each of the following sections four compounds are listed. In the box under each formula enter a number (1 to 4) indicating the order of acid strength (1 is strongest & 4 is weakest).

(a)	CH ₃ CH ₂ CH ₂ CO ₂ H	CH ₃ CH ₂ CHBrCO ₂ H	CICH2CH2CH2CO2H	CH₃CCl₂CO₂H
(4)	C.H.CH.OH	C ₆ H ₅ CO ₂ H	C ₆ H ₅ OCH ₃	0 11 011
(b)	C ₆ H ₅ CH ₂ OH	G611500 211	0611500113	C ₆ H ₅ OH
(c)	ОН	CO ₂ H		○
	NH ₂	N-H		
(d)			$N-CH_3$	O N-H

- 10. In the two questions below, you are asked to rank the relative strengths of illustrated acids and bases. Use your knowledge of resonance and inductive to answer this.
- A. For the series of bases shown below, rank the set from strongest to weakest.

(i)	NH		N	$\bigcap_{N \atop H}$	Strongest
	(a)	(b)	(c)	(d)	Weakest
(ii)	OMe		NMe ₂		Strongest
	(a)	(b)	(c)	(d)	Weakest

B. For the series of acids shown below, rank the set from strongest to weakest.

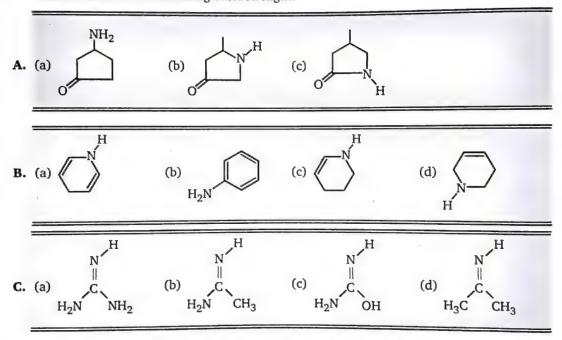
(i)	НО	но	HO NO ₂	HOCN	Strongest
	(a)	(b)	(c)	(d)	Weakest
(ii)	HO ₂ C OMe	HO ₂ C NO ₂	HO ₂ C	HO ₂ C NO ₂	Strongest
	(a)	(b)	(c)	(d)	Weakest

 In each of the following sections four compounds are listed. (Decreasing order of acidic strength, 1 is strongest & 4 is weakest).

(a)	CH ₂ (CO ₂ C ₂ H ₅) ₂	CH ₃ COCH ₂ CO ₂ C ₂ H ₅	(CH ₃ CO) ₂ CH ₂	RC == CH
(b)	RCH ₂ NO ₂	RSO ₂ CH ₃	(C ₆ H ₅) ₃ CH	RCOCH ₃
	$CH_2(C \equiv N)_2$	CH ₂ (NO ₂) ₂	HC ≡ N	RCH ₂ CO ₂ C ₂ H ₅
(c)				

(d) CH₂; CH₂; CH₂

12. Rank in the order of increasing basic strength.



13. Compare acidic strength of the following (Write your answer in box).

A.	OH	OH	OH	
	(a)	(b)	(c)	
В.	H ₂ N OH	H ₃ CO OH	F ₃ C OH	
	(a)	(b)	(c)	

GENERAL ORGANIC CHEMISTRY

41.

C.	CI OH	Cl FOH	F OH Br	·	
	(a)	(b)	(c)		
D.					
	(a)	(b)	(c)	(d)	
E.					
	(a)	(b)	(c)		
F.	(H)	H	(H)		
	(a)	(b)	(c) .		

14. Arrange the hydrogens in increasing order of their acidic strengths.

A.
$$H_a$$

B. H_c
 H_b
 H_b
 H_b
 H_d
 H

15. The compounds whose structures are shown below, incorporate a variety of functional groups. The question on the right ask you to identify which compounds have a specific functional group. For each compound that has the designed group, enter the appropriate number. The aromatic rings should not be counted as double bonds.

A.	Which have carbon-carbon double bonds ?
В.	Which have a ketone carbonyl group?
C.	Which have an aldehyde carbonyl group ?
D.	Which have aromatic rings ?
E.	Which have a hydroxy group ?
F.	Which have ether groups ?
G.	Which have an ester group ?
н.	Which have an amide group ?
I.	Which have a carboxylic acid group ?

(a) A

16.

Problem	A	В	С	D
1	O	CO ₂ H	O O	CH₂OH
2	ОН	OCH ₃	ОН	O ₂ N OH
3	N-H	N−H	N-H	$N-CH_3$
4	H C C C C CO ₂ H	H CI CO ₂ H	H C C CO ₂ H	CI CC CI CI

A.	Which is the stronge	st acid in 1?		
	(a) A	(b) B	(c) C .	(d) D
В.	Which is weakest aci	id in 1?		
	(a) A	(b) B	(c) C	(d) D
C.	Which is the stronge	st acid in 2?		
	(a) A	(b) B	(c) C	(d) D
D.	Which is weakest aci	id in 2?		
			() ((1)

- (c) C (d) D (b) B (a) A **E.** Which is the strongest acid in 3?
- (c) C (d) D (b) B (a) A
- F. Which is weakest acid in 3? (c) C (d) D
- **G.** Which is the strongest acid in 4?
- (d) D (b) B (c) C (a) A
- H. Which is weakest acid in 4? (c) C (d) D (b) B

17. For each of the six structural formulae (A through F), shown below, five questions are posed. The answer to each is a number that should be entered in the appropriate answer box.

The allower to each is a nur	mber mat should be entered in	the appropriate and
H ₃ C CH ₂	C≡C-CO ₂ H	N Z = Z
A	В	С
N−H S C≡N	H ₂ C=CH-CN-CH ₃ H ₃ C	CH ₃ OH CH ₃
D .	E	F

A.	(i) Number of sp^3 carbons : B.	Number of sp^3 C.	Number of sp ³
	(ii) Number of sp^2 carbons :	carbons:	carbons : Number of sp^2 carbons :
	(iii) Number of sp carbons:	Number of sp carbons:	Number of sp carbons:
	(iv) Number of carbon-carbonσ-bonds:(v) Number of π-bonds to	Number of carbon-carbon σ -bonds :	Number of carbon-carbon σ -bonds :
D.	carbon:	carbon:	Number of sp^3 carbons:
	(ii) Number of sp ² carbons:	Number of sp ² carbons:	Number of sp^2 carbons:
	(iii) Number of sp carbons:	Number of sp carbons:	Number of sp carbons:
	(iv) Number of carbon-carbon σ-bonds :	Number of carbon-carbon σ-bonds:	Number of carbon-carbon σ-bonds :
	(v) Number of π-bonds to carbon:	Number of π -bonds to carbon :	Number of π -bonds to carbon :

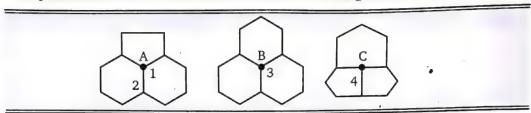
18. Match the column (I) and (II). (Matrix)

	Column (I)		Column (II)
	Molecule		Property
(a)		(p)	cis-compound
(ь)		(q)	trans-compound
(c)		(r)	Highest heat of combustion
(d)	\sim	(s)	lowest heat of combustion

19. Match the column (I) and (II).

. Mate	Column (I)		Column (II)
	Molecule		pK _a of Conjugate acid
(a)	N O H	(p)	0.8
(b)	$\bigcap_{\substack{N\\ CH_2-CH_3}}$	(q)	5.33
(c)	N N	(r)	10.65
(d)	N N O	(s)	10.95

20. The junctures centred on atoms A, B and C on the given structure.



A. Which juncture has the greatest deviation from planarity?

(a) A

(b) B

(c) C

(d) Cannot be predicted

B. Of the carbon-carbon bonds, (shown above) numbered from 1 to 4, which represent the most favourable site for H₂ addition?

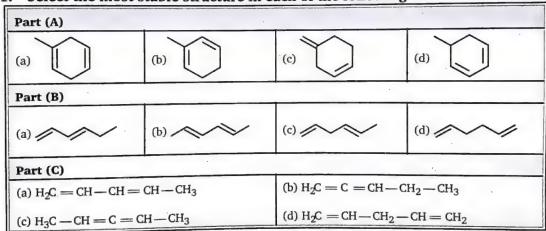
(a) 1

(b) 2

(c) 3

d) 4

21. Select the most stable structure in each of the following



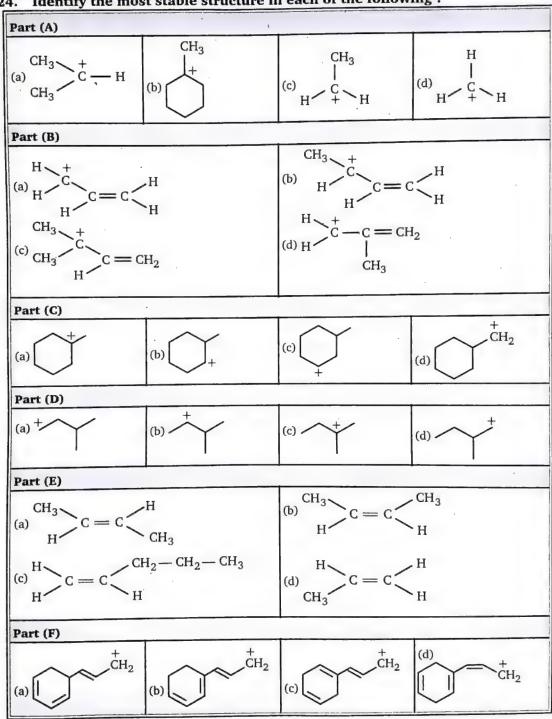
22. Match the column I and II. (Matrix)

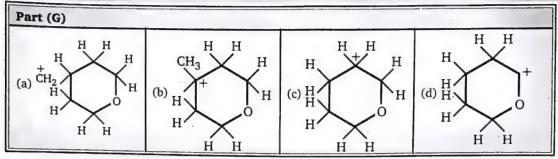
	Column (I)		Column (II)
(a)	-NO ₂	(p)	- m effect
(b)	-0-	(q)	+ m effect
(c)	-0-CH ₃	(r)	+ I effect
(d)	-C≡N	(s)	- I effect

23. Match the column I and II. (Matrix)

Column (I)		Column (II)
$H_3C - CH = CH - CH_3$	(p)	Dipole (cis > trans)
211 (21)	(g)	Dipole (trans > cis)
CII CI	(r)	Melting point ((trans > cis)
(c) $H_3C - CH = CH - CI$ (d) $CI - CH = CH - CI$	(s)	Boiling point (cis > trans)

24. Identify the most stable structure in each of the following:

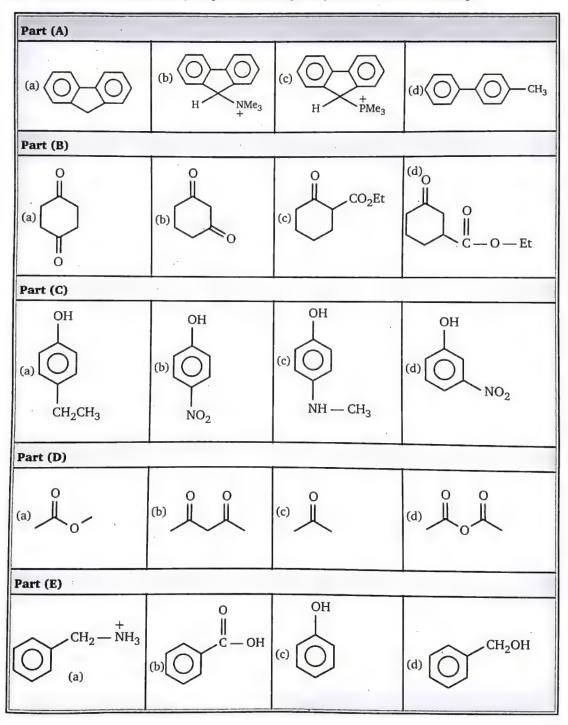




25. Identify the most basic compound in the following

	ost basic compound i		
Part (A)			
(a) NH	(b) NH	(c) NH	NH NH
Part (B)			
(a) N	(b) NH ₂	(c) NH ₂	$(d) \qquad \qquad NH - CH_3$
Part (C)			
(a) NaOH	(b) NaOOCCH ₃	(c) NaOCH ₃	(d) NaNH ₂
Part (D)			
(a) NH ₂	(b) NH ₂ NO ₂	(c) OCH ₃	(d) NH ₂

26. Identify the most acidic hydrogen containing compound from the following.



No. of			interition y just		
Part (F)	Part (F)				
(a) CH ₃ CH ₂ OH	(b) CH ₃ CH ₂ NH ₂	(c) $CH_3 - C \equiv CH$	(d) $CH_3 - CH = CH_2$		
Part (G)					
(a) CH ₃ — CO ₂ H	(b) CH ₂ —CO ₂ H NH ₃ ⊕	(c) NH ₃	(d) H ₃		
Part (H)					
OH NO ₂ NO ₂	(b) OH NO ₂	OH NO ₂ NO ₂	(d) NO ₂		
Part (I)					
CH CO	(b)	CH (c)	(d)		

27.

$$N \equiv C^{1} - C^{2} \Big|_{CH_{2} - C^{3} - H}$$

Give the type of hybridization present at each atom.

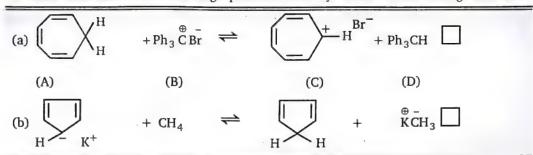
- (i) N —.....
- (ii) C₁ —.....
- (iii)C₂ —..... (vi)C₃ —.....

- (iv) O₁ —.....
- (v) CH₂ —.....

GENERAL ORGANIC CHEMISTRY

51

28. Predict the direction of the following equilibrium. Write your answer in the box given below.



29. Match the column I and II. (Matrix)

	Column (I)	•	Column (II)
(a)	NaHCO ₃ will react with	(p)	O OH OH Squaric acid
(b)	Na will react with	(q)	С-0-Н
(c)	NaOH will react with	(r)	О ОН
(d)	NaNH ₂ will react with	(s)	О 0 1 1 1 1 1 1 1 1 1

30. Match the column I and II.

	Column (I)	Column (II)	
	Acid	pK _a	
(a)	CH ₃ — CO ₂ H	(p) 5.69	
(b)	(CH ₃) ₃ NCH ₂ CO ₂ H	(q) 4.27	
(c)	⊕ (CH ₃) ₃ N(CH ₂) ₄ CO ₂ H	(r) 1.83	
(d),	O ₂ C — CH ₂ — CO ₂ H	(s) 4.80	

31. Match the column I and II.

	Column (I)		Column (II)
(a)	$\bigcirc \bigcirc $	(p)	NH ₃
(b)	$ \begin{array}{c} O \\ \parallel \\ C - O - H + NaHCO_3 \longrightarrow \end{array} $	(q)	14 CO ₂
(c)	$\bigcirc \bigcirc $	(r)	.co₂
(d)	$ \begin{array}{c} 0 \\ \parallel \\ S - O - H + NaNH_2 \longrightarrow \\ 0 \end{array} $	(s)	H ₂

SO₃H
OH
OH
$$CH - OH$$

$$C$$

Sum of molecular mass of gas (A + C) is:

Sum of molecular mass of gas A + B + C + D is :

34. Match the column I and II.

	Column (I)		Column (II)	
	Molecule	Rotatio	nal free energy barrier	
(a)	Ph	(p)	180 kJ/mol	
(b)	Ph P	(q)	88.3 kJ/mol	
(e)	Cl Cl Cl Ph	(r)	21 kJ/mol	
(d)	H C C H	(s)	Negative barrier	

35. Consider the following reaction of boron trifluoride (BF₃) and acetone:

$$BF_3$$
 + CH_3 \longrightarrow Product

- A. What is the critical HOMO (nucleophile) of this reaction?
 - (a) non-bonding orbital on boron
- (b) σ-orbital of acetone

(c) π -orbital of acetone

- (d) non-bonding electron pair orbital on oxygen
- **B.** What is the critical LUMO (electrophile) of the reaction?
 - (a) p-orbital of BF3

- (b) σ-orbital of BF₃
- (c) π^* orbital of acetone
- (d) non-bonding electron pair orbital on oxygen
- **C.** Which of the following is the correct product of this reaction?

(Lone electron pairs are not shown explicitly).

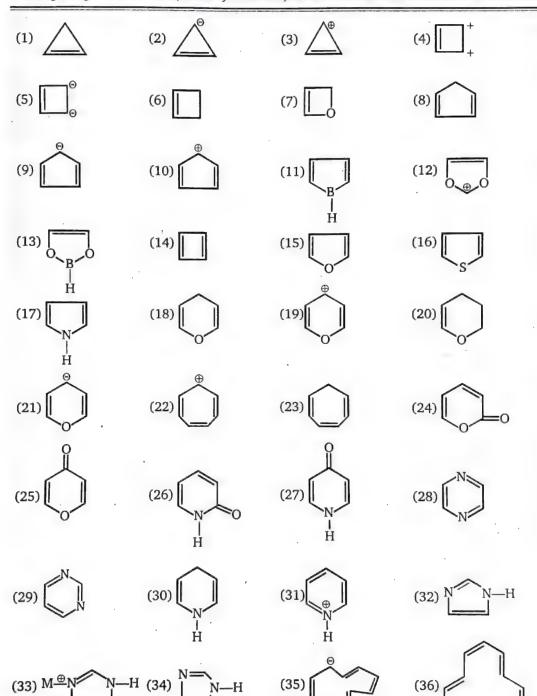
(a)
$$CH_3$$
 (b) $OOODD$ (c) $OOODD$ O

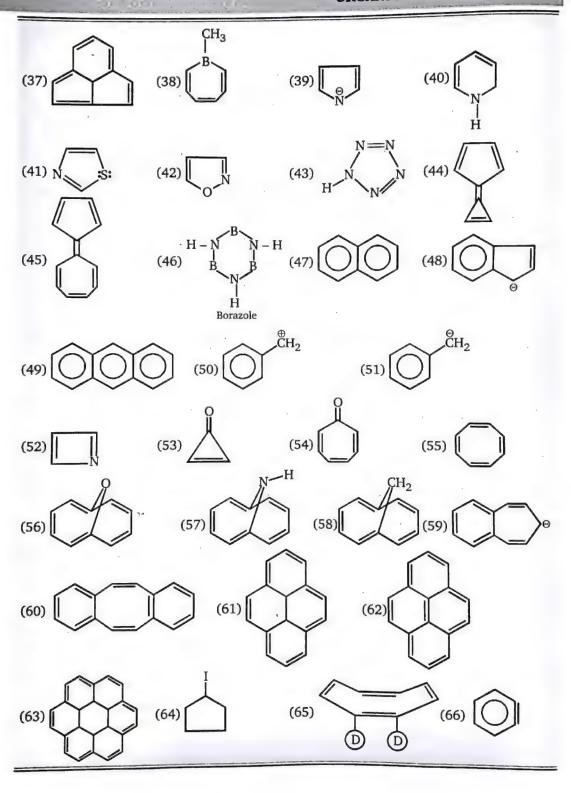
36. Rank the following carbocations according to stability (1 = most stable, 5 = least stable).

Put the answer in the boxes.



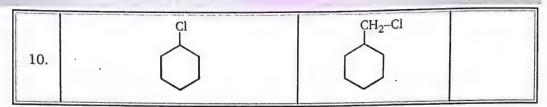
37. Among the given molecules, identify aromatic, anti-aromatic and non-aromatic molecules.





38. Among the given pairs, which is more reactive towards AgNO₃ (or) toward hydrolysis.

	Compound (A)	Compound (B)	Put the Answer here
1.	OBr	OBr	
2.	Br	Br	
3.			
4.	Br	Br	
5.	+ Cl	C1	
6.	CH ₃ -O -CH ₂ -Cl	CH ₃ -CH ₂ -CH ₂ -Cl	
7.	Br	CH ₂ -Br	
8.	Br	Br	
9.	Br	Br	



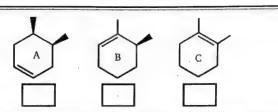
39. Put the answer in boxes given as directed.

S.No.	Property	Molecules	Correct Answer	Name of force responsi ble for the property
A.	highest boiling point	NCl ₃ ClNH ₂ NH ₄ Cl NH ₃		
В.	highest boiling point	☐ CH ₃		
C.	most soluble in water	OH OH OH SH		
D.	highest solubility in benzene	NH NH		

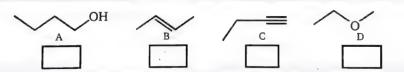
40. Circle any conjugated portions of these molecules.

$$CH_3O$$
 (A)
 $CH_2 = C = CH - CH = CH_2$
 (B)
 (C)
 (D)

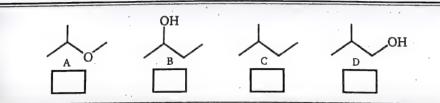
- 41. Arrange in the order as directed -
- A. The given alkenes in the order of their stability (1- most stable, 3-least stable).



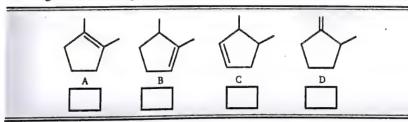
B. Arrange the following in the order of their acidic strength (1-most acidic, 4-least acidic)



C. Arrange the following molecules in order of expected boiling point. (1=highest bpt ; 4=lowest bpt.)



D. Arrange the following alkenes in order of their stability. (1 = most stable; 5 = least stable).



42. Match the column. (Matrix)

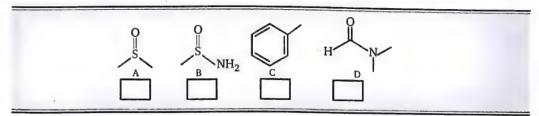
	Column (I)		Column (II)		
Compounds		Number of Benzylic hydrogen			
(a)		(p)	2		
(b)	CH ₂ -CH ₃ CH ₃	(q)	3		
(c)		(r)	4		
(d)		(s)	5		

43. Identify (+M) mesomeric & (-M) group of following.

	+M	-M	-I	+1
N				
\bigcirc				
NO				

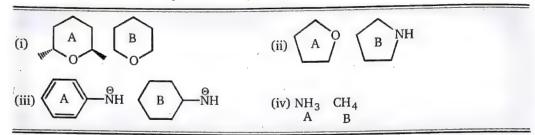
GEIVERUME OROMIVIC CITE		and the second second	
C - NH - CH ₃			
C – CH ₃			
0			
NO ₂			
OCH ₃			
CH = O			
O = C - OH			
NH - CH ₃			·

44. Identify the following solvents as polar protic (PP), polar aprotic (PA), non-polar protic (NPP) or non-polar aprotic (NPA).



62

45. Identify the stronger nucleophile in each pair.



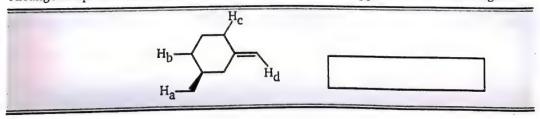
- 46. Encircle the molecule as directed:
 - (a) Which has higher boiling point: HBr or HCl
 - (b) Which has a higher boiling point : $CH_3 CH_2 OH$ or $CH_3 CH = O$
 - (c) Which is more miscible with methanol (CH₃OH): CH₃ CH₂ CH₂ CH₂ CH₃

or
$$CH_3 - CH_2 - CH_2 - CH_2 - CH_2 - CH_3$$

- (d) Which has a higher melting point: CH₄ or CH₃ CH₂ CH₃
- (e) Which has a higher boiling point: $CH_3 CH_2 CH_2 CH_2 CH_3$ or $CH_3 CH_2 CH_3$ 47. Encircle the molecule as directed:
 - (a) Which is more stable: BH₃ or BF₃
 - (b) Which is a stronger base: HO or H₂O
 - (c) Which is a stronger base: HO or HS
 - (d) Which is a stronger acid: HCl or HI
 - (e) Which is a stronger acid: HOCl or HCl
- 48. Explain why A has lower boiling point than B?

$$F_3C-CF_2$$
 F
 CF_3
 (A)
 (B)
 (B)
 (B)
 (CF_3)
 (DF_3)
 (DF_3)

49. Arrange the protons shown in the decreasing order of their approximate bond energies.



GENERAL ORGANIC CHEMISTRY

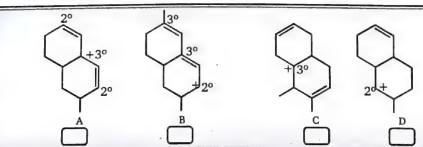
63

50. Consider the H-atoms in the molecule given below and answer the following.

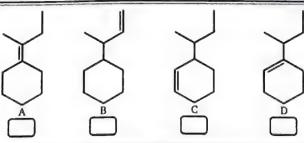
- (A) Identify the type (1°, 2° or 3° alkyl, vinyl, allyl etc.) of these H-atoms.
- (B) Arrange them in the decreasing order of their case of abstraction (easiest first)
- **51.** Consider the molecule shown below and answer with respect to $H_a \longrightarrow H_e$

$$H_a$$
 H_b
 H_c

- (A) Identify the type of H-atom (1°, 2°, 3° alkyl, vinyl or allyl)
- (B) Arrange them in decreasing order of their bond energy.
- **52**. Rank the following carbocations in order of stability (1 = most stable).



53. Rank the following alkenes according to energy (1 = lowest energy).



54. Match the column:

C11	Column (I)		Column (II)	
(Compounds)		(Double bond equivalent value)		
(a)		(p)	11	
(b)		(q)	12	
(c)		(r)	13	
(d)		(s)	14	
		(t)	15	

SUBJECTIVE PROBLEMS

1. How many 2° carbon in the following?

2. Find out the double bond equivalent (DBE) value of the given following compound:

$$\bigcup_{O} \bigcup_{NH_2}$$

3. Total number of functional groups present in the given following compound :

4. Total number of α -hydrogen in the given following compound is:



5. How many carbon atom present in the parent chain in the given following compound?

6. Total number of DBE value in :

- 7. How many isomers of $C_4H_{10}O$ reacts with Na metal to evolve H_2 gas ? (excluding stereoisomer)
- 8.

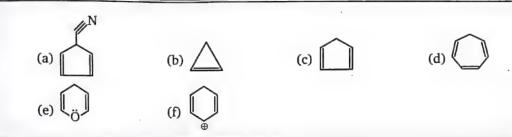
[x]-crown-[y]-ether.

value of
$$\frac{x+y}{3} = ?$$

9. Which of the given following compound will react with NaHCO3 or soluble in NaHCO3?

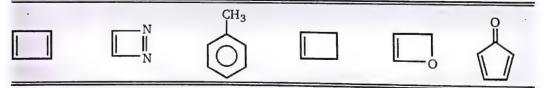
(e)
$$CH_3 - C \equiv CH$$

10. How many compound are stable after deprotonation?



Sum of types of functional group and DBE value for given compound is X so the value of X-10 is

12. P = Number of anti-aromatic compound, so the value of x is :



 $Q = \text{Total number of resonating structures of carbonate ion} [CO_3^{2-}]$

R =Number of α -hydrogen in given carbocation

S = Total number of geometrical isomers of CH $_3$ — CH = CH — CH = CH $_2$ T = Number of compound more acidic then CH $_3$ CH $_2$ OH

OH OH OH OH OH_3COOH , CH_3SO_3H

Sum of (P + Q + R + S + T) - 15 is:

13. x = number of (+M) group attached with phenyl ring, so the value of x is.

ANSWERS - LEVEL 2

```
1. a-4; b-3; c-2; d-1
```

2.
$$a-3$$
; $b-2$; $c-1$

3.
$$A - b$$
; $B - e$; $C - a$; $D - b$

4.
$$A - a$$
, c, f, g, k, l; $B - b$, d, h, j; $C - e$, i

5.
$$A - d; B - a$$

6.
$$C < B < A$$

8.
$$a - iv > ii > i > iii$$
; $b - iii > iv > i > ii$

9.
$$a-4$$
, 2, 3, 1; $b-3$, 1, 4, 2; $c-2$, 1, 3, 4; $d-2$, 3, 4, 1

10. A (i)
$$-d > b > c > a$$
; (ii) $-c > a > b > d$
B (i) $-c > d > b > a$; (ii) $-b > d > c > a$

12.
$$A-c < a < b$$
; $B-b < a < c < d$; $C-d < b < c < a$

13.
$$A-c < a < b$$
; $B-a < b < c$; $C-c < b < a$; $D-d < c < a < b$; $E-c < a < b$; $F-a > b > c$

14.
$$A - H_c < H_a < H_b$$
; $B - H_d < H_c < H_b < H_a$

16.
$$A - b$$
; $B - a$; $C - d$; $D - b$; $E - a$; $F - d$; $G - d$; $H - a$

18.
$$a-q$$
; $b-p$, r ; $c-p$, s ; $d-q$

19.
$$a - p$$
; $b - r$; $c - s$; $d - q$

20.
$$A - c$$
; $B - d$

21.
$$A - b$$
; $B - b$; $C - a$

22.
$$a - p$$
, s; $b - q$, r; $c - q$, s; $d - p$, s

23.
$$a-p, r, s; b-q, r; c-q, r; d-p, r, s$$

25.
$$A-d$$
; $B-a$; $C-d$; $D-c$

26.
$$A-c$$
; $B-b$; $C-b$; $D-b$; $E-b$; $F-a$; $G-b$; $H-c$; $I-b$

27. i.
$$-sp$$
; ii $-sp$; iii $-sp^2$; iv $-sp^2$; v $-sp^3$; vi $-sp^3$

29.
$$a-p. q, s; b-p, q, r, s; c-p, q, r, s; d-p, q, r, s$$

30.
$$a-s$$
; $b-r$; $c-q$; $d-p$

31.
$$a-r$$
; $b-q$; $c-s$; $d-p$

34.
$$a-q$$
; $b-r$; $c-s$; $d-p$

35.
$$A - d$$
; $B - a$; $C - a$

36.
$$A-3$$
; $B-1$; $C-5$; $D-2$; $E-4$

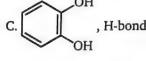
37. Aromatic—3, 4, 5, 9, 12, 13, 15, 16, 17, 19, 22, 24, 25, 26, 27, 28, 29, 31, 32, 33, 34, 35, 37, 38, 39, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 53, 54, 56, 57, 58, 61, 62, 63, 66

Non-aromatic-1, 6, 7, 8, 18, 20, 23, 30, 40, 55, 64, 65

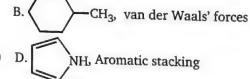
Anti-aromatic— 2, 10, 11, 14, 21, 36, 52, 59, 60

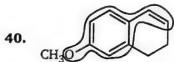
38.
$$1-B$$
; $2-A$; $3-B$; $4-A$; $5-A$; $6-A$; $7-B$; $8-B$; $9-A$; $10-A$

39. A. NH₄Cl, cation-anion interaction



, H-bonding (Also dipole-dipole)





$$CH_2 = C = CH - CH = CH$$





C.
$$A - 3$$
, $B - 1$, $C - 4$, $D - 2$

42.
$$a-s, b-r, c-q, d-p$$

B.
$$A-1$$
, $B-3$, $C-2$, $D-4$

D.
$$A-1$$
, $B-2$, $C-3$, $D-4$

43.

 $+\mathbf{M}$

-M

-I

`+I

(1) N

×

~

×

(2) NO

×

~

×

C - NH - CH₃

(3)

×

~

~

×

(4) C - CH₃

×

~

~

×

NO₂

×

~

~

×

OCH₃

~

×

~

×

CH = O

(7)

×

~

_

×

O = C - OH

(8)

×

~

~

×



- 44. A PA; B PP; C NPA; D PA
- **45.** (i) B, (ii) B, (iii) B, (iv) A
- **46.** (a) (HBr); (b) $(CH_3 CH_2 OH)$; (c) $(CH_3 CH_2 CH_2 CH_2 CH_3 CH_2 CH_3 CH_2 CH_3 CH_2 CH_3 CH_2 CH_3 CH_3 CH_3 CH_2 CH_3 -$
- 47. (a) (BF_3) ; (b) (HO); (c) (HO); (d) (HI); (e) (HOCI)
- **48.** In A, highly electronegative F-atoms are present at the periphery. In liquid term these F-atoms will repel each other due to partial negative charge and thus A will have lower b.pt.
- **49.** $H_d > H_a > H_b > H_c$
- **50.** A- H_a =1° alkyl; H_b = 2° alkyl; H_c =2° allyl; H_d = vinyl; H_e = 3° allyl B- Easiest to abstract: $H_e > H_c > H_b > H_a > H_d$ Hardest to abstract
- **51.** A- $H_a = 2^\circ$ alkyl; $H_b = 2^\circ$ allyl; $H_c = \text{vinyl}$; $H_d = 3^\circ$ allyl; $H_e = 1^\circ$ alkyl B- $H_c > H_e > H_a > H_b > H_d$
- **52.** A-2; B-1; C-3; D-4

53. A-1; B-4; C-3; D-2

54. a-r; b-t; c-t; d-s

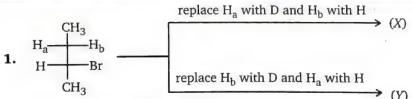
Subjective Problems

- **1.** 21 **2.** 11 **3.** 3 **4.** 6 **5.** 4 **6.** 13 **7.** 4 **8.** 7
- **9.** 2 (c, d) **10.** 3 (a, c, f) **11.** 7
- **12.** P = 3, Q = 3, R = 7, S = 2, T = 4 = 19 15 =**4 13.** 4

ISOMERISM (Structural & Stereoisomerism)



LEVEL-][



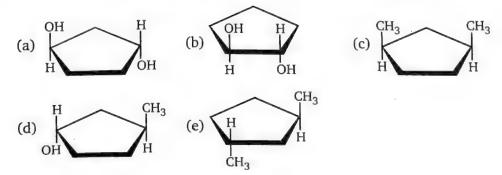
Relation between (X) and (Y) is:

(a) enantiomers

(b) diastereomers

(c) E and Z isomer

- (d) constitutional isomer
- 2. Which of the following cyclopentane derivative is optically inactive?



(d) 4

(d) 3

3. Which is the most stable conformer along the 2, 3 C - C bond axis of the compound?

4. Assign double bond configurations to the following:

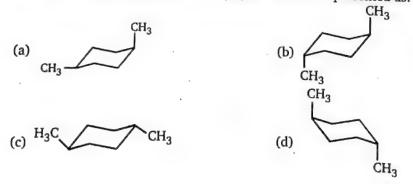
5. Allegra, a common prescription drug with the structure shown below, is given for the treatment of seasonal allergies. How many stereogenic carbon does Allegra possess?

6. How many meso isomers of C₄H₈Cl₂ will be?

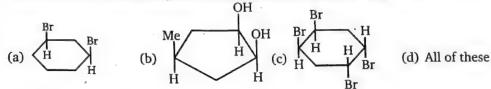
(a) 1

(a) 0 (b) 1 (c) 2

7. The stable form of trans-1, 4-dimethylcyclohexane is represented as:



8. Which of the following compound is non-resovable (meso) compounds?



9. $HO - CH_2 - CH_2 - F$

Which conformer of above compound is most stable across $C_2 - C_3$?

(a) staggered

(b) eclipsed (partially)

(c) gauche

- (d) fully eclipsed
- **10.** The following molecule is fluorometholone, a steroidal anti-inflammatory agent. How many stereogenic centers does it contain?

$$\begin{array}{c} \text{HO} \\ \text{H}_3\text{C} \\ \text{H} \\ \text{H} \\ \text{OH} \\ \text{H} \\ \text{OH} \\ \text{H} \\ \text{OH} \\ \text{OH}$$

(a) 5

(b) 6

(c) 7

(d) 8

11. How many chiral carbons are there in Reserpine (an antipsychotic drug)?

(a) 9

(b) 8

(c) 7

(d) 6

12. How many chiral centers are in the following compound?

(a) 4

(b) 5

(c) 6

(d) 7

13. Among the following, the optically inactive compound is:



14.
$$(C) \longrightarrow CH_3$$

$$(C) \longrightarrow CH_3$$

$$(D) \longrightarrow COOH$$

$$(D) \longrightarrow COOH$$

$$(C) \longrightarrow CH_3$$

$$(D) \longrightarrow COOH$$

$$(D) \longrightarrow COOH$$

$$(C) \longrightarrow CH_3$$

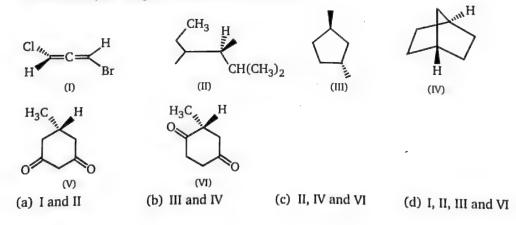
$$(C) \longrightarrow CH_3$$

$$(D) \longrightarrow COOH$$

15. Which of the following compounds might be useful to the chemist trying to increase the optical purity of the (d) sample?

(a)
$$HO_2C$$
 (b) HO_2C (c) CO_2H (d) CO_2H

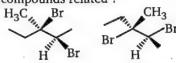
16. Which of the following molecules is (are) chiral?



17. The structure of (S)-2-fluorobutane is best represented by :



18. How are the following compounds related?



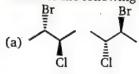
- (a) Diastereomers
- (c) Meso compounds

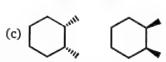
- (b) Enantiomers
- (d) Identical
- 19. Which one of the following is chiral?
 - (a) 1, 1-Dibromo-1-chloropropane
- (b) 1, 3-Dibromo-1-chloropropane
- (c) 1, 1-Dibromo-3-chloropropane
- (d) 1, 3-Dibromo-2-chloropropane
- 20. Among the following, the Newmann projections of meso-2, 3-butanediol are:

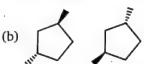
21. The binaphthol (Bnp) is:

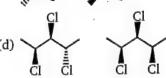
- (a) an optically active compound having chiral centre
- (b) an optically inactive compound
- (c) a meso compound
- (d) an optically active compound without having chiral centre

22. Which of the following pairs of compounds is a pair of enantiomers?









23. The maximum number of stereoisomers that could exist for the compound below?

$$\bigvee_{Br}^{Br} \bigvee_{Br}^{Br}$$

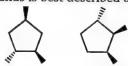
(a) 6

(b) 8

(c) 10

(d) 16

24. The following pair of compounds is best described as:



(a) identical

(b) diastereomers

(c) enantiomers

(d) none of the above

25. Determine the absolute configurations of the labeled carbons (a and b):



(a) a = R; b = R

(b) a = R; b = S

(c) a = S; b = R

(d) a = S; b = S

26. Which of the structures (a – d) will be produced if a "ring flip" occurs in the following compound in chair form?



(a) **(**







27. Which of the following compounds is most stable?



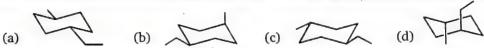






28. Which is the most stable chair form of this compound?



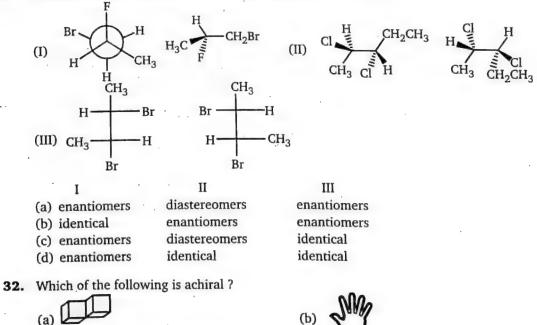


29. Which pairs of the salts would have identical solubilities in methanol?

(I)
$$H \xrightarrow{Ph} CH_3 CH_3 \xrightarrow{Ph} CO_2$$
 (II) $H \xrightarrow{NH_3^{\oplus}} CH_3 \xrightarrow{Ph} COO^-$
(III) $NH_3^{\oplus} CH_3 H \xrightarrow{Ph} CH_3 (IV) \xrightarrow{Ph} H CH_3 \xrightarrow{Ph} CH_3 H$
(IV) $Ph \xrightarrow{NH_3^{\oplus}} CH_3 H$

30. The following compounds differ in respect of :

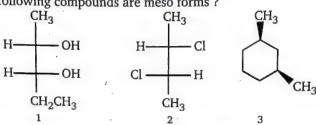
- (a) their chemical and physical properties
- (b) nothing
- (c) the direction in which they rotate plane of polarized light
- (d) their interactions with molecules
- 31. Indicate whether each of the following pairs are identical, or?





(d) a molecule of 3-methylheptane

33. Which of the following compounds are meso forms?

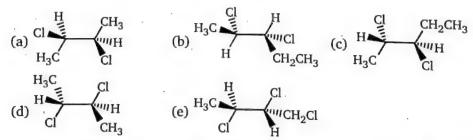


- (a) 1 only
- (c) 1 and 2

- (b) 3 only
- (d) 2 and 3
- **34.** The separation of a racemic mixture into pure enantiomers is termed as:
 - (a) Racemization
- (b) Isomerization
- (c) Resolution
- (d) Equilibration
- 35. Rank of the following groups in order of R, S precedence (IV is highest):

- CH(CH	₃) ₂	– CH₂CH₂Br		- CH ₂ Br		- C(CH ₃) ₃	
1		2		3		4	
I	П	III	IV	I	\mathbf{II}	III	ΙV
(a) 3	2 .	4	1	(b) 1	4	2	3
(c) 3	4	1	2	(d) 3	4	2	1

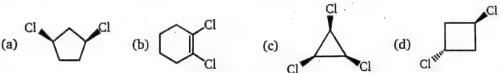
36. Which of the following is a meso compound?



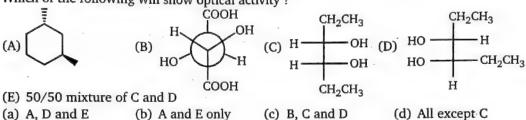
37. Among the following structures, select E isomers (arrows indicate the bonds to be considered)?

$$H_3C$$
 CH_3
 H_3C
 CH_3
 H_3C
 CH_3
 H_3C
 CH_3
 CH_3

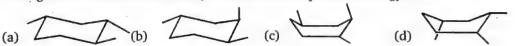
38. Which of the following compounds has a zero dipole moment?



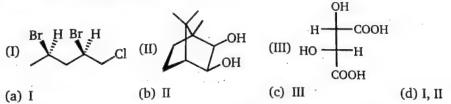
- **39.** On Pluto, where everything is frozen, astronauts discovered two forms of butane gauche and anti. Assuming that there are no rotations around single bonds, which statement about the two forms is correct?
 - (a) They are enantiomers
 - (b) They are diastereoisomers
 - (c) They are meso compounds
 - (d) The gauche form has two stereogenic centers, and the anti has only one
- 40. Which of the following will show optical activity?



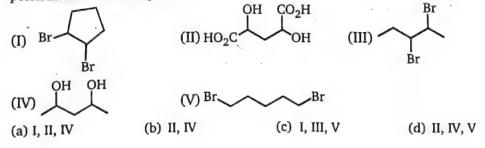
41. Among the structure shown below, which has lowest potential energy?



42. Which of the following molecules is/are chiral?

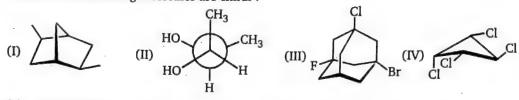


43. A compound was synthesized by a student, but its structure was not identified. However, his wonderfully helpful instructor told him that it was a meso compound with 5 carbons and 2 stereogenic centers. Which of the following structures should the student consider as possibilities for his compound?



44. How many isomers are possible for the following molecule?

45. Which of the following molecules are chiral?

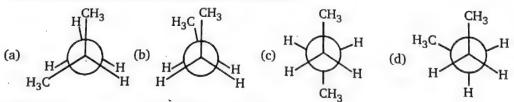


(a) I, II, III and IV

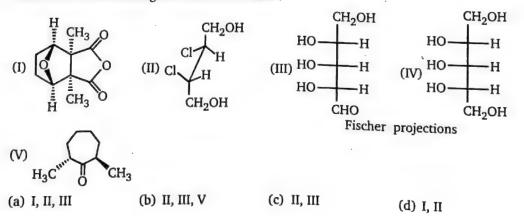
(a) 1

- (b) II, III and IV
- (c) II and IV
- (d) I and II
- **46.** Which equilibrium is not rapid at room temperature?

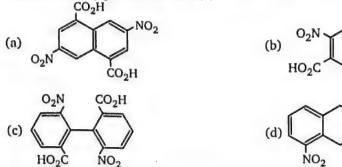
47. Which is the lowest energy conformation of butane?



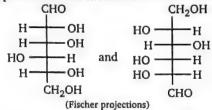
48. Which of the structures given below are chiral?



49. Which of the following carboxylic acids could be resolved by reaction with an enantiomerically pure chiral amine?



50. What is the relationship between the molecules in the following pairs?



(a) enantiomers

(b) diastereomers

(c) identical

(d) structural isomers

51. What are the correct designations for the structure below?

(a) E, E

(b) Z, E

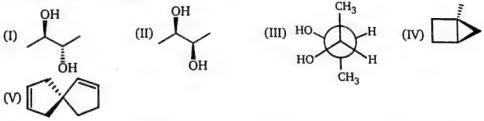
(c) E, Z

(d) No geometrical isomers are possible

CO₂H

CO₂H

52. Which of the following molecules are chiral?



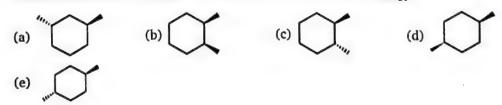
(a) I and III

(b) I and V

(c) II and III

(d) II, III, IV

53. Which one of the following isomeric structures has the lowest energy?



54. The following compounds are identical with respect to:

- (a) molecular composition
- (c) melting point

- (b) boiling point
- (d) IUPAC name
- **55.** Among the following, the most stable isomer is :
 - (a) OCH₃
 - (c) OMe OH

- (b) OH OMe
- (d) OH OMe
- **56.** The most stable conformation of the following compound is:

- (a) t-Bu Me
- (b) t-Bu Me
- (c) t-Bu H Me
- (d) t-Bu Me
- 57. Which of the following molecules have non-zero dipole moments?
 - (I) gauche conformation of 1, 2-dibromoethane
 - (II) anti conformation of 1, 2-dibromoethane
 - (III) trans-1, 4-dibromocyclohexane
- (IV) cis-1, 4-dibromocyclohexane

(V) tetrabromomethane

(VI) 1, 1-dibromocyclohexane

(a) I and II

(b) I and IV

(c) II and V

(d) I, IV and VI

58. What is the maximum number of stereoisomers possible for discodermolide?

(a)
$$2^{14}$$
 (b) 2^{15} (c) 2^{16} (d) 2^{17}

- **59.** An aqueous solution containing compounds A and B shows optical activity. A and B are stereoisomers. Which of the following possibilities cannot be correct?
 - (a) A has two chiral centers, but B does not have any because it has a symmetry plane
 - (b) A and B are enantiomers
 - (c) A and B are diastereomers
 - (d) A and B are not present in equal amounts
- **60.** Which of the following structures represents the lowest-energy form of (1S, 2S, 4R)-trimethyl-cyclohexane?

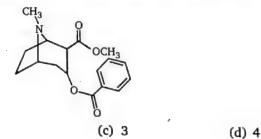


- 61. Which one of the following is a diastereomer of (R)-4-bromo-cis-2-hexene?
 - (a) (S)-4-bromo-cis-2-hexene
 - (b) (S)-5-bromo-trans-2-hexene
 - (c) (R)-4-bromo-trans-2-hexene
 - (d) (R)-5-bromo-trans-2-hexene

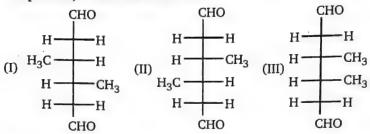
(a) 1

(b) 2

62. The structural formula of cocaine is shown below. How many stereogenic carbon atoms are there in this molecule?



63. Which of the following statements best describes the stereochemical relationships of compound I, II and III shown below?



- (a) All compounds are chiral
- (b) None of the compounds is chiral
- (c) I and II are meso compounds
- (d) I and II are diastereomers, and III is a meso compound
- (e) I and II are chiral
- **64.** What is the absolute configuration of the following molecules? (NS = the molecule has no center) Note: For the purpose of this question only, the order of stereocenters is not specified; *i.e.*, R, S = S, R.

(III)
$$H_2G=N$$
 $CH=NH$

- I II III IV
 (a) R R, S R NS
 (c) R R, S NS NS
- I II III IV
 (b) R R, R S R, R
 (d) R R, S R R, S
- **65.** The number of all the possible stereoisomers formed by the given compound is :

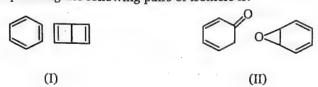
(a) 2

(b) 3

(c) 32

(d) 64

66. The relationship among the following pairs of isomers is:



I	A: Constitutional			
п	B: Configurational			
Ш	C: Conformational			
IV	D: Optical			

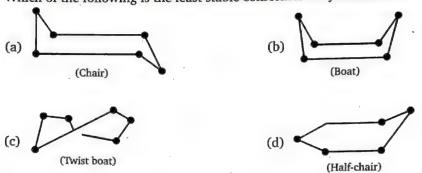
- (a) I-A, II-B, III-B, IV-D (c) I-B, II-A, III-B, IV-D
- (b) I-A, II-A, III-B, IV-D
- (d) I-B, II-B, III-A, IV-B
- The structural formula of sativene is shown below. How many stereogenic centers are there 67. in this molecule?



- (a) 2
- (c) 4

- (b) 3
- (d) 5

68. Which of the following is the least stable conformer of cyclohexane?



69. The S- enantiomer of ibuprofen is responsible for its pain-relieving properties. Which one of the following structures shown below is (S)-ibuprofen?

(c)
$$C = OH$$
 (d) $C = OH$ C

70. Which of the following depict the same?

- (a) 1 and 2
- (c) 2 and 3

- (b) 1 and 3
- (d) 1, 2, and 3

71. A naturally occurring substance has the constitution shown below. How many may have this constitution?

HO
$$CH_2OH$$
HO $CH = CHCH = CHCH_2CH_2CH_3$
(b) 8 (c) 16 (d) 64

72. The absolute configurations of the two centers in the following molecule are :



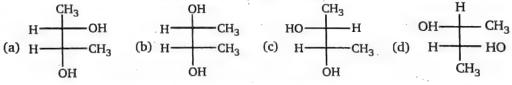
- (a) 2(R), 3(S)
- (b) 2(R), 3(R)
- (c) 2(S), 3(S)
- (d) 2(S), 3(R)
- 73. The total number of stereoisomer possible for 2, 3-dichloro butane :
 - (a) 2

(a) 2

(b) 3

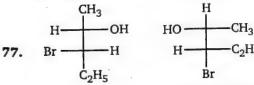
(c) 4

- (d) 5
- 74. Which of the following structure is not meso-2,3-butanediol?



- **75.** A solution of optically active 1-phenylethanol racemizes in acidified aqueous medium. It is due to:
 - (a) enolization

- (b) carbonium ion formation
- (c) carbanion formation
- ' (d) reversible oxidation-reduction
- 76. The most stable conformation of ethylene glycol is:
 - (a) Anti
- (b) Gauche
- (c) Partially eclipsed (d) Fully eclipsed



The molecules represented by the above two structures are :

(a) identical

(b) enantiomers

(c) diastereomers

- (d) epimers
- 78. The correct order of priority of groups $-SCH_3$ (I), $-NO_2$ (II), -C = CH (III) and $-CH_2C_6H_5$ (IV), on the basis of CIP classification, is (increasing order):
 - (a) I, III, II, IV

(b) IV, III, II, I

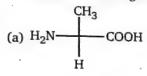
(c) II, IV, I, III

(d) III, IV, II, I

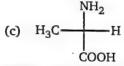
79. The configuration at C-2 and C-3 of the compound given:



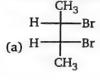
- (a) 2R, 3S
- (b) 2S, 3R
- (c) 2S, 3S
- (d) 2R, 3R
- 80. Amongst the following amino acids, the (R) enantiomer is represented by:



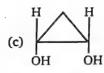
(b) H NH₂



- (d) H_3C \longrightarrow NH_2
- 81. Which of the following is a meso compound?



$$CH_2 - CH_3$$
 $H \longrightarrow OH$
 $CH_2 - CH_3$
 $CH_2 - CH_3$

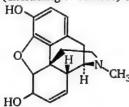


- (d) All of these
- **82.** Predict stereochemistry of product when d and l-amine reacts with ℓ -acid:
 - (a) Diastereomers

(b) Meso

(c) Racemic

- (d) Pure Enantiomer
- 83. How many chiral center (excluding N centres) are there in morphine?



(a) 4

(b) 5

(c) 6

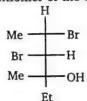
- (d) More than 6
- 84. Which dimethylcyclobutane is optically active?
 - (a) trans-1, 2

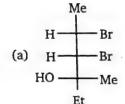
(b) cis-1, 2

(c) trans-1, 3

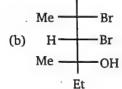
(d) cis-1, 3

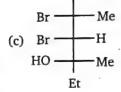
Which of the following is the enantiomer of the compound shown below?

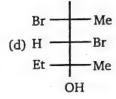




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86. How many different stereoisomers are possible for the following compound?

$$CIHC = HC - C - CH = CHCI$$

$$CI$$

- (a) 1
- (b) 2
- (c) 3
- (d) 4
- The following compounds are best described as: 87.
 - (R)-PhCH(OH)CH₃ and (S)-PhCH(OH)CH₃
 - (a) enantiomers
 - (b) diastereomers
 - (c) not stereoisomers
 - (d) conformational isomers (differing by single bond rotation)
- Rank the following substituent groups in order of decreasing priority according to the 88. Cahn-Ingold-Prelog system:

$$-CH(CH_3)_2$$
 $-CH_2Br$ $-CH_2CH_2Br$
 $1 > 3 > 1$ (b) $1 > 3 > 2$ (c) $3 > 1 > 3 > 2$

- (a) 2 > 3 > 1

- (d) 2 > 1 > 3
- Compare the stabilities of the following two compounds: 89.

A: cis-1-Ethyl-3-methylcyclohexane

- B: trans-1-Ethyl-3-methylcyclohexane
- (a) A is more stable

(b) A and B are of equal stability

(c) B is more stable

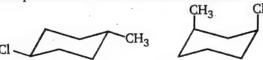
- (d) No comparison can be made
- What, if anything, can be said about the magnitude of the equilibrium constant K for the 90. following equilibrium?

$$H$$
 $CH(CH_3)_2$
 H
 $CH(CH_3)_2$
 H
 $CH(CH_3)_2$

- (a) K = 1
- (c) K > 1

- (b) K < 1
- (d) No estimate of K can be made

91. What is the relationship between the two structures shown?

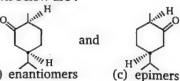


- (a) Constitutional isomers
- (b) Stereoisomers
- (c) Different drawing of the same conformation of the same compound
- (d) Different conformation of the same compound
- 92. Which of the following statements is true?
 - (a) van der Waals' strain in cis-1, 2-dimethylcyclopropane is the principal reason for its decreased stability relative to the trans isomer
 - (b) Cyclohexane gives off more heat per CH2 group on being burned in air than any other cycloalkane
 - (c) The principal source of strain in the boat conformation of cyclohexane is angle strain
 - (d) The principal source of strain in the gauche conformation of butane is torsional strain
- Ph CH = $NO_2H \xrightarrow{\text{isomerises}} (x)$, Isomer (x) is: 93.
 - (a) $Ph NO CH_2OH$

(b) Ph — CH₂ — NO₂

(c) $Ph - NH - CO_2H$

- (d) None
- Which of the following will not show geometrical isomerism? 94.
 - (a) $CH_3 C = CH CH_2 CH_3$
- (b) $CH_3 CH CH = CH CH_2 CH_3$
- (c) $CH_3 CH = CH CH_3$
- (d) $CH_3 CH_2 CH = CH CH_2 CH_3$
- The two compounds shown below are: 95.



- (a) diastereomers
- (b) enantiomers

(d) regiomers

The molecular formula of diphenylmethane, 96.

$$CH_2$$
 , is $C_{13}H_{12}$;

How many structural isomers are possible when one of the hydrogen is replaced by a chlorine atom?

- (b) 4
- (c) 8
- (d) 7
- Correct configuration of the following molecule is: 97.



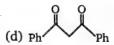
- (a) 2S, 3S
- (b) 2S, 3R
- (c) 2R, 3S
- (d) 2R, 3R

98. Maximum enol content is in :









99. Which of the following will have one of the stereoisomer meso?

(a) 2-chlorobutane

- (b) 2, 3-dichlorobutane
- (c) 2,3-dichloropentane

(d) 2-hydroxypropanoic acid

100. The correct decreasing order in the enol content of following molecules is:







(a) I > II > III

(b) II > I > III

(c) III > II > I

(d) II > III > I

101. Total number of stereoisomers of the compound 1-bromo-3-chlorocyclobutane is:

(a) 0

(b) 1

(c) 2

(d) 3

102. Total number of stereoisomers of the 1,3-dichlorocyclohexane is:

(a) (

(b) 1

(c) 3

(d) 4

103. Total number of stereoisomers of the compound 1, 4-dichlorocyclohexane is :

(a) 0

(b) 1

(c) 2

(d) 4

104. Total number of stereoisomers of the compound 2-4-dichloroheptane is:

(a) 0

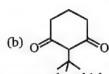
(b) 2

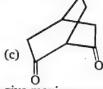
(c) 3

(d) 4

105. In which of the following keto form is more dominating than enol form:



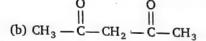




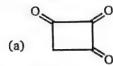
(d) all of these

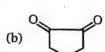
106. Among the following compounds, which will give maximum enol content in solution:

(a)
$$C_6H_5 - C - CH_2 - C - C_6H_5$$



107. Which of the following has unstable enol form?



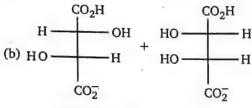


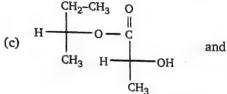


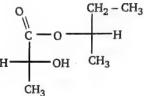


108. Calculate enantiomeric excees of mixture containing 6g of (+) 2-butanol and 4g of (-) -2-butanol.

- (a) 10%
- (b) 20%
- (c) 40%
- (d) 33%
- 109. Which of the following pair represent pair of diastereomers?
 - (a) Meso tartaric acid and (l) tartaric acid

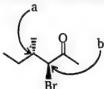






- (d) All of these
- **110.** The stereochemistry of this molecule is :

- (a) 1R, 3R
- (b) 1R, 3S
- (c) 1S, 3S
- (d) 1S, 3R
- 111. Pure (S)-2-butanol has a specific rotation of +13.52 degrees. A sample of 2-butanol prepared in the lab and purified by distillation has a calculated specific rotation of +6.76 degrees. What can you conclude about the composition?
 - (a) 50% (S), 50% impurity
- (b) 50% (S), 50% (R)
- (c) 50% (S), 50% racemic
- (d) some other mixture
- 112. Determine the absolute configurations of the chiral centres in the following compound.



(a) a = R; b = S

(b) a = R; b = R

(c) a = S; b = S

- (d) a = S; b = R
- 113. Total number of stereoisomers possible for following compound is:

$$\begin{array}{c} \text{CH} = \text{CH} - \text{CH}_2\text{CH}_3 \\ \text{CH} = \text{CH}_2 \end{array}$$

- (a) 8
- (b) 16
- (c) 32
- (d) 64

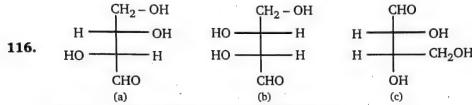
Which is the correct structure of D-glyceraldehyde?

(a)
$$H \xrightarrow{CHO}$$
 OH (b) $HO \xrightarrow{H}$ CHO (c) $HO \xrightarrow{CH_2OH}$ (d) All of these CH_2OH OHO

115.
$$HO - CH_2 - CH_2 - CH_2 - CH_1 - H$$

Which conformer of above compound is most stable (consider conformer across (C₂ - C₃)

- (a) Staggered
- (b) Gauche
- (c) Fully eclipsed
- (d) Partially eclipsed

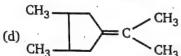


- (D) & (L) Configuration of above carbohydrate is:
- (a) L, L, D (b) L, D, L
- (c) L, L, L
- (d) L, D, D
- How many isomers have the name bromomethylcyclopentane? (ignoring chirality) 117.
- (b) 5
- (c) 6
- Which of the following compound can show geometrical isomerism? 118.

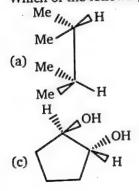
(a)
$$\frac{Br}{I} = C = C \frac{Cl}{Cl}$$

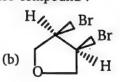
(p)
$$C < CH^3$$

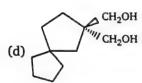
(c)
$$\sum_{Cl}^{F} C = C \sum_{Et}^{Et}$$



119. Which of the following structure represent meso-compound?







(d) both (b) & (c)

How many representations of lactic acid are possible in Fischer projection (d & l)?

(a) 8

(b) 12

(c) 24

(d) 36

121. Total number of stereoisomer formed by the given compound is:

(a) 2

(b) 3

(c) 4

(d) 8

122. The number of stereoisomers formed by the given compound is:

(a) 2

(b) 3

(c) 4

124.

(d) 5

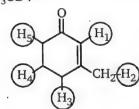
123. Which of the following compound does not undergo base - catalyzed exchange in D_2O even though it has an α -hydrogen?

(a) (b) (c) OH
$$\frac{D_2O/OH^-}{long time}$$
 Product

Identify the product formed in the above reaction:

(d) None of these

125. In 3-methyl-2-cyclohexenone which hydrogen cannot undergo deuterium exchange when it reacts with CH $_3$ O $^{\Theta}$ /CH $_3$ OD?



(a) H_1, H_4

(c) H_3, H_2

- (b) H_4
- (d) H_5, H_3

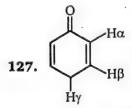
126. (II)



The tautomer of II is:

- (c) both I and III

- (b) III
- (d) none of these



In the enolization of the given molecule, the H-atom involved is:

(a) α-H

(b) β-H

(c) γ-H

(d) cannot be enolized

Among the given structure which can exhibit tautomerism?

(a) I only (b) II only CH_3 129. CH_3

 CH_3 (II)

(d) none of these (c) III only CH₃ CH_3 CH₃ CH_3 (III)

Identify the which can exhibit tautomerism?

(a) I only

(b) II only

(c) III only

(d) all of these

130.
$$CH_3 - CH = O \longrightarrow CH_2 = CH - OH$$
(I) (II)

Between the two tautomers which is more stable?

(c) I = II (b) II (a) I 131.

(d) none of these

Correct stability order of the given tautomers is:

(a) I > II > III

(b) III > II > I

(c) II > I > III

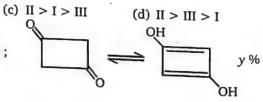
z %

(d) II > III > I

HO. 132. (II)

Correct stability order of the given tautomers is:

(a) I > II > IIII < II < III (d) 133.



The correct order of enol contents x, y, z is:

(a)
$$x > y > z$$

(b)
$$z > y > x$$

(c)
$$y > x >$$

(d)
$$x > z > y$$

z%(x,y,z represent enol content)

The correct order of x, y, z is:

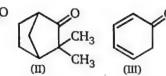
(a)
$$x > y > z$$

(b)
$$z > y > x$$

(c)
$$y > x > z$$

(d)
$$x > z > y$$

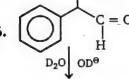




Among the given ketones, the one which does not enolize is:

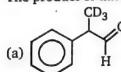
- (a) I
- (b) II
- (c) III
- (d) none of these

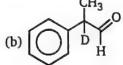


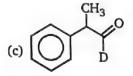


Product

The product of this reaction should be:







(d) All of these





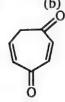


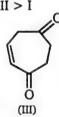
Among the given compounds, the correct order of enol content is :

- (a) I > II > III
- I < II < III (d)
- (c) II > I > III
- $(d) \parallel > \parallel \parallel > 1$

138.







Among the given compounds, the correct order of enol content is:

- (a) I > II > III
- (b) III > II > I
- (c) II > I > III
- I < III > II

(II)



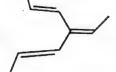
Among the given compounds, the correct order of enol content is:

(a) I > II > III

(I)

- I < II < III (d)
- (c) III > I > II
- III < I < II (b)

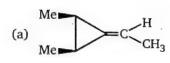
140.

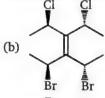


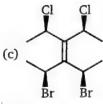
How many geometrical isomers are possible for the above compound?

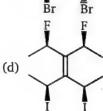
- (b) 4
- (c) 6
- (d) 8

141. Which of the following compound will not show geometrical isomerism across the π -bond?









Choose the correct relation between l_1 and l_2 ?

- (a) $l_1 = l_2$

- (b) $l_1 > l_2$ (c) $l_1 < l_2$ (d) $l_2 = 2l_1$

143.

Choose the correct relation between l_1 and l_2 ?

- (a) $l_1 = l_2$
- (b) $l_1 > l_2$ (c) $l_1 < l_2$
- (d) $l_2 = 2l_1$

 CH_3 144.

How many geometrical isomers are possible for the above compound?

- (a) 0
- (b) 2
- (c) 3
- (d) 4

 CH_3 145. C_2H_5

How many geometrical isomers are possible for the above compound ?

(a) 0

(b) 2

(c) 3

(d) 4

ĊH3 146. CH_3

How many geometrical isomers are possible for the above compound?

(a) 0

(b) 2

(c) 3

(d) 4

 CH_3 147.

Br How many geometrical isomers are possible for the above compound?

(a) 0

(b) 2

(d) 4

 CH_3 148.

How many geometrical isomers are possible for the above compound?

(a) 0

(b) 2

(c) 3

(d) 4

 CH_3 CH₃ 149.

How many geometrical isomers are possible for the above compound?

(d) 4

150.

I and II are geometrical isomers of each other because

(a) $l_1 = l_2$

(b) $l_1 > l_2$

(c) $l_2 > l_1$

(d) l_1 and l_2 cannot be compared.

151. $CH_2 = CH - CH = CH - CH = CH_2$

How many geometrical isomers are possible for this compound?

(a) 2

(b) 3

(c) 4

(d) 8

152.
$$CH_3 - CH = C - C = CH - CH_3$$

How many geometrical isomers are possible for this compound?

(b) 3 (d) 6

- (c) 4
- **153.** $CH_3 CH = C C = CH - CH_3$

How many geometrical isomers of this compound are possible ?

(a) 2

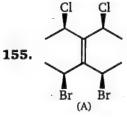
(b) 3

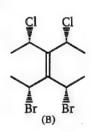
(c) 4

(d) 6

- (a) chiral
- (c) Optically active

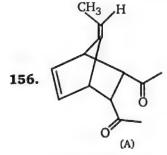
- (b) C₃ axis of symmetry
- (d) All of these

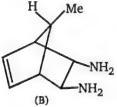




Relationship between above pair (A) & (B) is:

- (a) Enantiomer
- (b) Diastereomers
- (c) Identical
- (d) Structural isomer





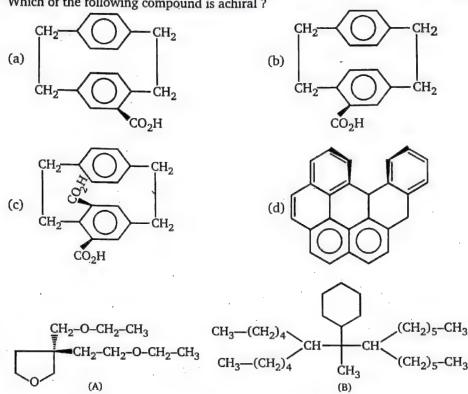




From the above compound (A), (B), (C) & (D) chiral compound is:

- (a) A
- (c) C
- (d) D

157. Which of the following compound is achiral?



R and S configuration of compound (A) & (B) will be:

(a) R, R (c) S, R

158.

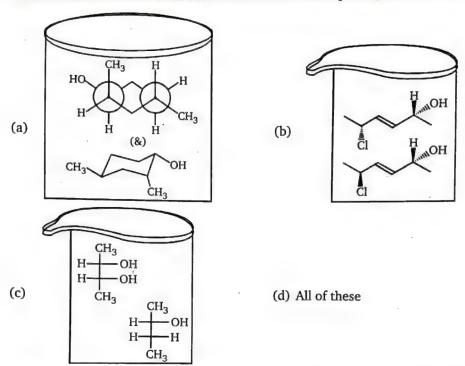
(b) R, S

(d) S, S

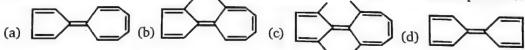
159. Which of following compound has center of symmetry?

(a)
$$CH_2$$
 CH_2 CH_3 $CH_$

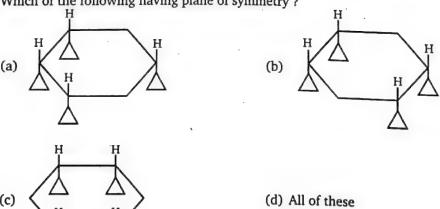
160. Which mixture of structure in each beaker would rotate plane polarized light?



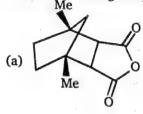
161. Which of following compound will rotate the plane polarized light at room temperature?

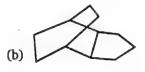


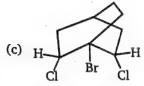
162. Which of the following having plane of symmetry?



163. Which of following compound is achiral?







R = -CH - Cl

(d) All of these

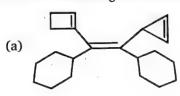
164. Which of the following compound has plane of symmetry?

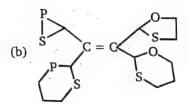
S = -CH - Cl Br

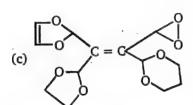


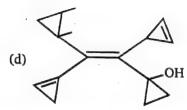
(d) None of these

165. Which of following is E isomer?

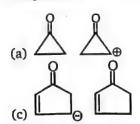


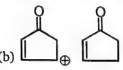






166. Among the given pairs, in which pair second compound has less enol content than first compound?





(d) none of these

167. Which of the following is incorrect relation between given pairs?

(d) 6

168. (B); (A) and (B) are isomer and isomerization effectively OH (A)

carried out by trace of base (B). Identify (B).

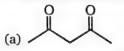
(a)
$$Ph - CH_2 - C - O - H$$
 (b) $Ph - C - O - CH_3$ (c) $Ph - C - CH_2 - OH$ (d) $H - C - CH_2 - O - Ph$

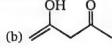
 ${\rm CH_3-CH=CH-CH=CH-CH_3};$ total number of geometrical isomer is : 169. (b) 3

(a) 2 (c) 4

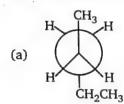
Identify most stable enol form of terric acid: 170.

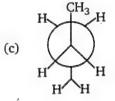
Which structure is most stable?

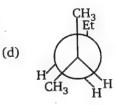




172. Identify conformer of 2-methly pentane:

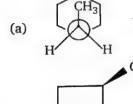




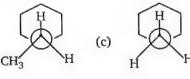


173. The lowest energy conformer of





(b)



(d)

174.

How many atoms will be bisect during plane of symmetry?

(b) 4

(c) 6

(d) 8

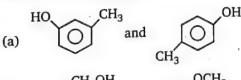
175. The number of all types of isomers of chlorobutane is:

(b) 4

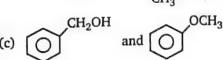
(c) 6

(d) 5

176. Which of the following pairs of compounds are not positional isomers?



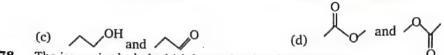
and



(d) All of these

177. Which of the following pairs of compounds are functional isomers?

(a)
$$OH$$
 and OO



178. The isomeric alcohol which has a chiral carbon atom is:

(a) n-butyl alcohol (b) iso-butyl alcohol (c) sec-butyl alcohol (d) tert-butyl alcohol

179. The pair of enantiomers among the following compound is:

(a) I and IV

(b) II and IV

(c) II and III

(d) I and II

180. Which of the following is chiral?

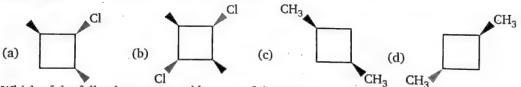
(a) Cell phone

(b) Spiral staircase

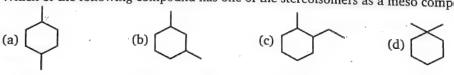
(c) Scissor

(d) All of these

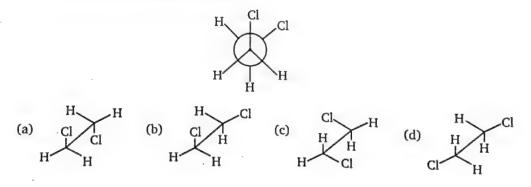
181. In which of the following compound, possess plane of symmetry as well as centre of symmetry?



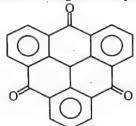
182. Which of the following compound has one of the stereoisomers as a meso compound?



183. For the following Newman projection

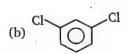


184. Which of the following is correct for the given compound?

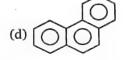


- (a) It possess centre of symmetry
- (b) It possess C₄ axis of symmetry
- (c) It possess plane of symmetry
- (d) Compound is chiral
- **185.** Which of the following molecules has axis of symmetry and a coaxial plane of symmetry?









- (e) All of these
- **186.** Number of diastereomer of given compound :



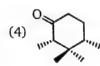
- (a) 2
- (b) 3
- (c) 4
- (d) 6
- **187.** Which of the structures is/are diastereomer of **A**?





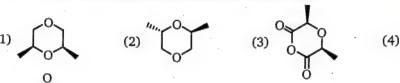




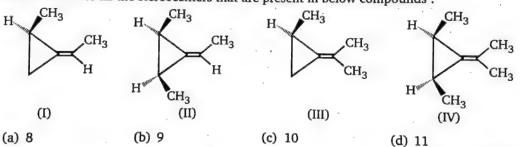


- (5)
- (a) 3
- (b) 1 and 4
- (c) 2 and 3
- (d) 5

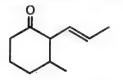
188. Identify which of the structures below are meso structures?



- (5)
- (a) 1 and 3
- (b) 1.3 and 5
- (c) 1,3 and 4
- (d) 2 and 5
- 189. How many enol form is possible for $CH_3 CH_2 CH_2 CH_2 CH_3$ (including stereoisomers) will be?
 - (a) 2
- (b) 3
- (c) 4
- (d) 5
- **190.** Find the sum of all the stereocenters that are present in below compounds:



- **191.** A pair of stereoisomers might be classified in various ways. Which of the following statement are true with respect to pairs of stereoisomers?
 - (a) They might be configurational isomers
- (b) They might be diastereomers
- (c) They might be constitutional isomers
- (d) They might be tautomers
- (e) They might be conformational isomers
- (f) They might be enantiomers
- (g) They might be positional isomers
- (a) a, b, c, e
- (b) b, d, e, f, g
- (c) a, b, f
- (d) a, b, c, f
- 192. Ignoring specific markings, which of the following objects are chiral?
 - (I) a shoe
- (II) a book
- (III) a pencil
- (IV) a pair of shoes (consider the pair as one object)
- (V) a pair of scissors
- (a) I only
- (b) I & V
- (c) I, IV, V
- (d) III, IV, V
- 193. Calculate the total number of stereoisomers when alkene having trans configuration:



- (a) 2
- (b) 3
- (c) 4
- (d) 8

194.
$$(A) \xrightarrow{D_2O/DO^-} (B)$$

After prolonged treatment of (A) by D_2O/DO^- , the difference in molecular weights of compounds (A) and (B) is:

(a) 2

(b) 3

(c) 4

(d) 8

195. $\xrightarrow{\text{Cl}_2}$ A mixture of all isomers possible from the mono-chlorination of the structure is subjected to fractional distillation, then how many fractions will be obtained?

(a) 2

(b) 3

(c) 4 ·

(d) 5

196. Number of optically active isomer is/are:

(a) 0

(b) 1

(c) 2

(d) 3

197. At normal temperature, *X* and *Y*

(a) resonance structures

(b) tautomers

(c) functional isomers

(d) positional isomers

198. Two possible stereoisomers for

(a) enantiomers

(b) diastereomers

(c) conformers

(d) rotamers

199. The configurations of the carbon atoms C₂ and C₃ in the following compound are respectively

(a) R, R

(b) S, S

(c) R, S

(d) S, R

200. The compound that is chiral is

(a) 3-methyl-3-hexene

(b) 4chloro-1-methycyclohexane

(c) 2-phenylpentane

(d) 1, 3-disopropylbenzene

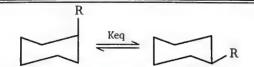
						ANICV	TEDS	- LEV	/FT 1						
1.	(b)	2.	(c)	3.	(b)	4.	(c)	5.	(a)	6.	(b)	7.	(c)	8.	(d)
9.	(c)	10.	(d)	11.	(b)	12.	(c)	13.	(a)	14.	(b)	15.	(b)	16.	(d)
17.	(c)	18.	(a)	19.	(b)	20.	(b)	21.	(d)	22.	(b)	23.	(c)	24.	(d)
25.	(a)	26.	(b)	27.	(d)	28.	(b)	29.	(a)	30.	(c)	31.	(c)	32.	(a)
33.	(b)	34.	(c)	35.	(c)	36.	(d)	37.	(c)	38.	(d)	39.	(b)	40.	(a)
41.	(a)	42.	(d)	43.	(a)	44.	(d)	45.	(a)	46.	(b)	47.	(c)	48.	(b)
49.	(c)	50.	(c)	51.	(d)	52.	(d)	53.	(e)	54.	(a)	55.	(d)	56.	(c)
57.	(d)	58.	(b)	59.	(a)	60.	(a)	61.	(c)	62.	(d)	63.	(e)	64.	(d)
65.	(b)	66.	(b)	67.	(d)	68.	(d)	69.	(d)	70.	(d)	71.	(d)	72.	(a) ·
73.	(b)	74.	(a)	75.	(b)	76.	(b)	77.	(a)	78.	(b)	79.	(c)	80.	(b)
81.	(d)	82.	(a)	83.	(b)	84.	(a)	85.	(a)	86.	(d)	87.	(a)	88.	(d)
89.	(a)	90.	(b)	91.	(a)	92.	(a)	93.	(b)	94.	(a)	95.	(b)	96.	(b)
97.	(a)	98.	(d)	99.	(b)	100.	(a)	101.	(c)	102.	(c)	103.	(c)	104.	(d)
105.	(d)	106.	(a)	107.	(c)	108.	(b)	109.	(d)	110.	(a)	111.	(c)	112.	(c)
113.	(a)	114.	(d)	115.	(b)	116.	(b)	117.	(c)	118.	(d)	119.	(b)	120.	(c)
121.	(a)	122.	(b)	123.	(d)	124.	(b)	125.	(b)	126.	(c)	127.	(c)	128.	(b)
129.	(d)	130.	(a)	131.	(c)	132.	(d)	133.	(d)	134.	(d)	135.	(b)	136.	(b)
137.	(c)	138.	(a)	139.	(d)	140.	(b)	141.	(b)	142.	(a)	143.	(c)	144.	(b)
145.	(b)	146.	(b)	147.	(b)	148.	(a)	149.	(b)	150.	(c)	151.	(a)	152.	(c)
153.	(b)	154.	(d)	155.	(c)	156.	(a)	157.	(c)	158.	(d)	159.	(d)	160.	(d)
161.	(b)	162.	(d)	163.	(d)	164.	(d)	165.	(d)	166.	(c)	167.	(d)	168.	(c)
169.	(b)	170.	(c)	171.	(c)	172.	(d)	173.	(b)	174.	(c)	175.	(d)	176.	(c)
177.	(b)	178.	(c)	179.	(c)	180.	(d)	181.	(d)	182.	(b)	183.	(b)	184.	(c)
185.	(e)	186.	(b)	187.	(b)	188.	(a)	189.	(c)	190.	(c)	191.	(c)	192.	(b)
193.	(c)	194.	(c)	195.	(b)	196.	(a)	197.	(b)	198.	(a)	199.	(a)	200.	(c)



LEVEL-2

1. Match the Column (I) and (II).

	Column (I)	Column (II)			
	Reaction		Stereoisomers		
(a)	$CH_3 - CH = CH - CH = N - OH$	(p)	2		
(ъ)		(q)	4		
(c)	$CH_3 - CH = CH - CH = CH - CH = CH - CH_3$	(r)	6		
(d)	$CH_3 - CH = CH - CH = CH - CH = CH - Ph$	(s)	8		



	Column (I)	Co	olumn (II)
	Group	Equilib	rium Constant
(a)	R = -H	(p)	38
(b)	$R = -CH_3$	(q)	23
(c)	R = -Et	(r)	18
(d)	$R == - CH - CH_3$ CH_3	(s)	1

	Column (I)		Column (II)	
6-	Molecule	Nature		
(a)	$\begin{array}{c} \overset{H}{=} \\ \overset{CO_2CH_2CH_2OH}{=} \\ \overset{E}{\to} \\ CO_2H \end{array}$	(p)	Chiral	
(b)	CO ₂ CH ₂ CH ₂ O ₂ C H	(q)	Achiral	
(c)	OH H CO ₂ H CO ₂ H OH	(r)	Meso	
(d)	HO CO ₂ H H	(s)	Compound containing even number of chiral centers	

. Ma	Column (I) and (II). (M	g 0	Column (II)			
	Compound	Isomerism				
(a)	CH ₃ H	(p)	Geometrical isomerism			
(b)	CH ₃ Et	(q)	Optical isomerism			
(c)	CH ₃ H Et	(r)	Compound containing plane of symmetry			
(d)	CH ₃ CH ₃	(s)	Compound containing center of symmetry			

5. Match the Column (I) and (II).

E	Column (I)		Column (II)
	Molecules		Relationship
(a)	Cl and CH ₃	(p)	Identical
(b)	Cl and Cl CH ₃	(q)	Enantiomer
(c)	Cl and CH ₃	(r)	Diastereomer
(d)	Cl and Cl CH ₃	(s)	Structural Isomerism

	Column (I)	A:	Column (II)
	Compound		Nature
(a)	CH_3 $OH CH_3$ CH_3 $OH CH_3$ CH_3	(p)	cis-compound
(b)	CH_3 $OH CH_3$ CH_3 CH_3	(p)	trans-compound
(c)	CH ₃	(r)	Optically active
(d)	CH ₃	(s)	Optically inactive

7. Match the Column (I) and (II). (Matrix)

Column (I)			Column (II)	
	Molecule	Property		
(a)	CI $C = C = C$ CI	(p)	Chiral centers containing compound	
(b)	CH_3 CH_3 H	(q)	Presence of stereocenter	
(c)	Br — F	(r)	Optically active compound	
(d)	CH_3 $C = N$ OH	(s)	. Compound containing plane of symmetry	

117

(many	Column (I)	municipated in space.	Column (II)		
	Molecule	Property			
(a)	F > C = C = C = C < F	(p)	Polar molecule		
(ь)	$rac{F}{H}$ $C = C = C < rac{H}{F}$	(p)	Optically active		
(c)	F > mil F	(r)	Optically inactive		
(d)	H H H H	(s)	Plane of symmetry		

	Column (I)		Column (II)			
	Molecule	Property				
(a)	Me V _{min} H	(p)	Meso compound			
(b)	Me VH Me Me	(p)	Achiral			
(c)	Me Vann H Inn H Me	(r)	Chiral compound			
(d)	Me Me Me	(s)	Compound will show geometrical isomerism			

ISOMERISM 119

	Column (I)		Column (II)	
M	odified Newmann Projection	Conformers		
(a)	CH ₃ CH ₃	(p)	Fully eclipsed	
(b)	CH ₃ H CH ₃	(4)	Partially eclipsed	
(c)	H CH ₃	(r)	Gauche	
(d)	H CH ₃ CH ₃	(s)	Staggered	

	Column (I)		Column (II)	
	Newmann Projection	Name of the Compound		
(a)	H CH ₃ CH ₃ CH ₃	(p)	3-methyl pentane	
(b)	CH ₃ CH ₃ H	(q)	n-butane	
(c)	CH ₃ H	(r)	Methyl-cyclopentane	
(d)	H CH_3 H CH_3 CH_3 H H	(s)	1,2,4-trimethyl cyclohexane	

12. Match the Column (I) and (II). (Matrix)

-	Column (I) Column (II)			
	Molecule '	Property		
(a)		(p)	Rotates plane polarized light	
(b)	Br	(p)	Cannot rotate plane polarized light	
(c)	Br AsMe ₃	(r)	Plane of symmetry	
(d)	CH_3 $C = C = C_{min}^{CH_3}$	(s)	Centre of symmetry	

G	Column (I)	Column (II)		
	Molecule		Stereocenters	
(a)	$CH_3 - CH = CH - CH - CH_3$ Br	(p)	1	
(b)	$H-C \equiv C-CH = CH-CH-CH-CH_3$ Br Br	(q)	2	
(c)	$\begin{array}{c} \text{O} \\ \parallel \\ \text{Ph} - \text{S} - \text{CH} = \text{CH} - \text{CH} - \text{CH}_3 \\ \parallel \\ \text{CH}_3 \end{array}$	(r)	3	
(d)	Ph — CH — Et	(s)	4	

	Column (I)		Column (II)	
	Molecule	Stereoisomers		
(a)	CCCCMe	(p)	2	
(b)	C C COMe	(q)	0	
(c)	C C COMe	(r)	4	
(d)	C C C COMe	(s)	8	

Ė	Column (I)	Column (II)		
	Molecule	Property		
(a)	Me C — Me	(p)	Meso Compound	
(b)	Me_2N Me OH $C = C - Me$	(q)	Compound having even no. of chiral centres	
(c)	O N H	(r)	Optically active compound	
(d)	СООН Н — ОН СООН	(s)	Compound having odd no. of chiral centres.	

16. Match the Column (I), (II) and (III). (Matrix)

	Column (I)		Column (II)		Column (III)
	Property		Molecule	No. of Chiral Cente	
(a)	CH_3 $C = C$ $CHDCI$	(p)	Optically active	(w)	0
(b)	CH ₃	(q)	Optically inactive	(x)	1
(c)	$\begin{array}{c} O^{\ominus} \\ \\ CH_3 \stackrel{\oplus}{} N {} HCl^{\ominus} \\ \\ Et \end{array}$	(r)	Plane of symmetry	(y)	2
(d)	Cl	(s)	Centre of symmetry	(z)	3

17.

(a)	H₃C CH₃	(b)	CH ₃ CH ₃	(c)	H
(d)	H $C = C = C$ H	(e)	CH_3 $C = C = C $ H	(f)	H CH3
(g)	H Br CH ₃ CH ₃ Br H	(h)	CH ₃ CH ₃		·

From the above compounds select :

(A)	two of which are chiral and contain chiral centre :	
(B)	two of which are achiral and contains chiral centre :	
(C)	two of which are chiral and does not contain chiral centre :	
(D)	two of which are achiral and does not contain chiral centre :	

18. Comprehension

Consider the given structures and answer A, B & C.

A. Which of the compound is optically active?

(a) P

(b) R

(c) S

(d) T

B. Which of the isomer is most stable?

(a) R

(b) S

(c) T

(d) U

19. Identify relationship between following pairs:

(a)	$\begin{array}{c ccccc} & & & & & & & & & & & \\ & & & & & & & $	(b)	Cl F Br Br H Cl
(c)	OH OH	(d)	H ₃ CH ₃ H ₃ CH ₃ CH ₃ F H ₄ CH ₃ CH ₃ F H CH ₃

If they are enantiomer answer will be 1, if they are diastereomers answer will be 2, if they are constitutional isomers answer will be 3 and if they are identical present 4 as the answer. Sum of answer of each part a + b + c + d is :

20. In each of the following three questions a hydrocarbon is named. For each select from among the sixteen conformational structures (a through p) all structures that represent possible conformers of that compound. Write letters (a through p), corresponding to your selections, in each answer box.

A.	2-methylbutane	
В.	2,3-dimethylpentane	
C.	1-ethyl-1, 3-dimethyl cyclohexane	

L	11-ethyl-1, 3-dimethyl cyclol	lexan	e		
(a)	H CH ₃ CH ₃ CH ₃ CH ₃ H	(b)	Н3С Н	(c)	H_3C H_3C H_3C CH_3
(d)	H ₃ C W CH ₃	(e)	H CH ₃	(f)	H CH ₃ C ₂ H ₅ CH ₃
(g)	H H_3C CH_3 H C_2H_5 CH_3	(h)	CH ₃ CH ₃ CH ₃	(i)	H ₃ C - H ₅ CH ₃
(j)	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	(k)	H_3C C_2H_5 CH_3	(1)	CH ₃ CH ₃ CCH ₅ CC ₂ H ₅
(m)	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	(n)	CH ₃ CC ₂ H ₅	(o)	H_3C CH_3 C_2H_5
(p)	$\begin{array}{cccccccccccccccccccccccccccccccccccc$		·		

21. Examine structures a through j, shown below, with respect to their symmetry or lack of it. Assume that the five-membered rings and the ring in compound g are planar. The wedge-hatched bonds in b, c, d & e designate specific configurations. Also, for the acyclic compounds assume stable anti conformations. Answer each of the following questions by writing letters (a through j), corresponding to your selections, in each answer box. If there is no structure that fits the description enter an x in the answer box.

Α.	Which structures are chiral ?	·
В.	Which structures have a plane of symmetry?	
C.	Which structures have a center of symmetry?	

(a)	Н	(b)	Br	(c)	Br
(d)	Br	(e)	Br min Br	(f)	OH
(g)		(h)	C ₂ H ₅ CHCl ₂	(i)	C₂H₅CHClC₂H₅
(j)	C ₂ H ₅ CHClCH ₃				

- 22. (i) 1,2-dichlorocyclopropane = w
 - (ii) 1,3-dimethyl-cyclobutane = x
 - (iii) 2-bromo-3-chlorobutane = y
 - (iv)1,3-dimethyl cyclohexane = z

Calculate total number of stereoisomer of the above compounds.

Sum of w + x + y + z =

23. Examine the following formulas and select those pairs that satisfy the following conditions: Be sure to write two letters (and only two) in each answer box, unless you select f. In the second and third parts more than one answer is possible.

(a)	H CH ₃ C ₂ H ₅	(b)	H_3 C H_3 C H_3	(c)	H ₃ C CH ₃
(d)	H ₃ CW, H	(e)	H ₃ C CH ₃ CH ₃ H	(f)	No formulas meet the designated condition

A.	Which are identical in all respects?	
В.	Which are conformational isomers?	
C.	Which are constitutional isomers?	

24. Examine the following formulas and select those pairs that satisfy the following conditions: Be sure to write two letters (and only two) in each answer box. In the second and fourth parts more than one answer is possible.

(a)	Br H	(b)	Br H Br	(c)	Br H Br
(d)	H Br Br H	(e)	H H Br	(f)	Br H Br

A.	Which are identical in all respects?	
В.	Which are configuration isomers?	
C.	Which are conformational isomers?	

25. Consider the following statements regarding the given projection (True or False).

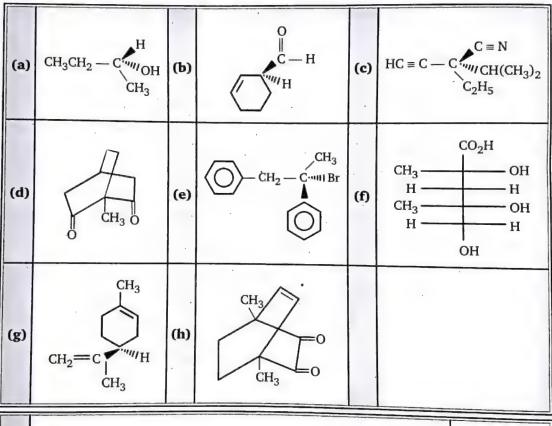
(w)	$\begin{array}{c c} \text{CH}_2\text{Cl} \\ \text{H} & \text{Br} \\ \text{Cl} & \text{H} \\ \text{CH}_3 \end{array}$	(x)	$\begin{array}{c} \text{CH}_2\text{Cl} \\ \text{Cl} & \text{H} \\ \text{H} & \text{Cl} \\ \text{CH}_3 \end{array}$
(x)	CH ₃ H CH ₂ Cl	(Z)	$\begin{array}{c} \text{CH}_2\text{Cl} \\ \text{H} & \text{Cl} \\ \text{CH}_3 \end{array}$

(a)	W and Y are diastereomers
	Z is the projection of X
(c)	W, X, Y and Z are optically active
	Y and Z are isomer

26. Examine the following structural formulas and select those that are chiral.

٠.	Examine the following structural formulas and select those that are third.							
	(a)	OH	(b)	NH ₂ CO ₂ H	(c)			
	(d)	ОН	(e)	$\begin{array}{c c} & CO_2H \\ H & OH \\ H & OH \\ \hline & CH_2-CO_2H \end{array}$	(f)	$H \longrightarrow OH$ $HO \longrightarrow CH_3$ $CH_2 - CO_2H$		
	(g)	CH3	(h)	O_2 O_3 O_3 O_3 O_3 O_3	(i) _.	CH_3 $C = C = C$ H		
	(I)	$H \xrightarrow{CH_3} CI$ $H \xrightarrow{CH_3} CI$ CH_3						
		Write your						

27. The configuration of eight compounds, a through h are shown below, using various kind of stereo representations. To answer the question given below, write (a through h) indicating your choice.



A.	Which of these configuration are achiral?
В.	Which configuration has no stereogenic center?
c.	Which configuration has more than one stereogenic center?
D.	Which of these configuration are meso compound?

28. The structural formula of ten compounds, (I) through (X) are drawn below, you may select any one of these structure.

Answer the following question about that compound.

I	CH ₃ /m.	п	CH ₃	ııı	HO CH ₃
IV	CH ₃	v	CH ₃	VI	CH ₃ <u>≡</u> CH ₃
VII	CH ₃	VIII	CH3 OH	ıx	CH ₃ /m, OH
x	CH ₃ OH				

A. How many chiral centre are present in this compound?

- (a) 0 (e) 4
- (b) 1 (f) 5
- (c) 2
- (d) 3

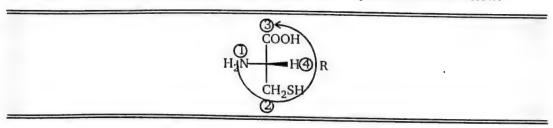
B. Is this compound chiral or achiral?

- (a) Chiral
- (b) Achiral

C. What symmetry element are present in this compound?

- (a) None
- (b) Plane of symmetry
- (c) Center of symmetry

29. The structure of one of the enantiomers of the amino acid cysteine is shown below.



Classify this structure as:

(a) R or S

(b) D or L

30. Identify the following double bonds either E, Z or None (N) in the compounds given below either.

A.

B. (a) Bongkrekic acid is a toxic compound produced by Pseudomonas cocovenenans, and isolated from a mold that rows on bongkrek, a fermented Indonesian coconut dish. (a) Label each double bond as E, Z or neither (N).

- (b) How many total stereoisomers (including all types) are possible for bongkrekic acid?
- (c) How many sites of unsaturation are present in bongkrekic acid?

ISOMERISM 133

31. Designate the following double bonds as E, Z or none (N) configuration in the boxes provided below.

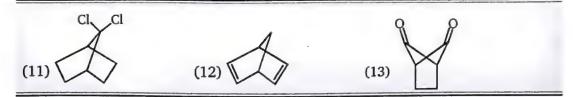
32. The following compounds may exist as two or more stereoisomers. These may be classified as enantiomer pairs or meso compounds.

Answer the following question about the above structure.

- (A) Total number of stereoisomers:
- (B) Number of enantiomeric pairs :
- (C) Number of meso compounds:

33. Identify axis of symmetry in the given compound.

ISOMERISM 135

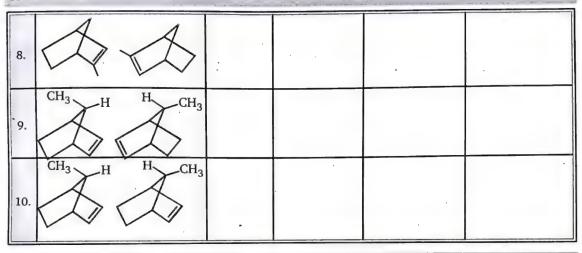


34. For each of the following pharmaceutical compounds, identify all stereogenic (i.e., all asymmetric carbon atoms) and label the configuration of each as being either (R) or (S).

35. Find relationship between given pair :

		Identical	Enantiomer	Diastereomer	Constitutional Isomer
1.	CO ₂ H CO ₂ H			·	
2.					
3.	OH OH				
4.	CH ₃ Et H OH HO H H OH HO H Et CH ₃				
5.	H CH ₃ CH ₃ H CH ₃				
6.	CH ₃ CH ₃ CH ₃ CH ₃ H H H				
7.	44				

137 **ISOMERISM**



36. Comprehension

Structural formula of compound (A) is following:

- The correct statement(s) about the compound (A) is/are:
 - (a) The total number of stereoisomers possible for (A) is 3
 - (b) The total number of mesoisomer possible for (A) is 1
 - (c) The total number of pair of enantiomer possible for (A) is 1
 - (d) All of these
- Number of plane of symmetry in cis-form of compound (A) is: В.
- (b) 1
- (c) 2
- (d) 3

37. Match the column. (Matrix)

	Column (I)		Column (II)	
-	No. of Carbon	No. of structural ison		
(a)	C ₄ H ₁₀	(p)	2	
(b)	C5H12	(q)	3	
(c)	C ₆ H ₁₄	(r)	5	
(d)	C ₇ H ₁₆	(s)	9	

38. Match the column. (Matrix)

	Column (I) Column (II)				
	Compound	% of enol content			
(a)	Ot.	(p)	100 %		
(ъ)		(q)	76 %		
(c)	$ \begin{array}{c c} O & O \\ & \\ CH_3 - C - CH_2 - C - CH_3 \end{array} $	(r)	8%		
(d)	O O CH ₃ - C - CH ₂ - C - O - Et	(s)	Keto-Enol is not possible		

39. Draw a most stable conformation (N - C) bond in the following compound.

$$H \stackrel{H}{\sim} C = C \stackrel{H}{\sim} H$$

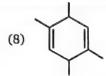
40. Find total number of stereoisomers for each compound given below:

O || (1)
$$Ph - S - CH = CH - CH_2 - CH = C = C = CH - CH = CH - CH_3$$

(2)
$$CH = CH - CH_3$$
 $NO_2 CI$

(3)
$$Cl$$
 CH_3
 CH_3

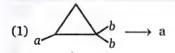
N-OH



(9)
$$\frac{\text{Cl}}{\text{H}}$$
 $\text{CH} - \text{CH} = \text{CH} - \text{CH}_3$

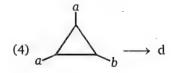
(10)
$$CH_3 - CH = CH - CH - CH_3$$

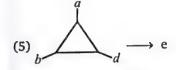
41. Find the total number of stereoisomer for each compound :











(6)
$$a \longrightarrow f$$

42. Match the column:

	Column (I)		Column (II)	
	Pair	Isomeric Relationship		
(a)	$CH_3 \longrightarrow C \longrightarrow CH_3$ $CH_3 \longrightarrow C \longrightarrow CH_3$	(p)	Chain	
(b)	$\begin{array}{c} \text{O} \\ \\ \text{CH}_3 - \text{CH}_2 - \text{CH}_2 - \text{C} - \text{OH} \text{, CH}_3 - \text{CH} - \text{CH}_3 \\ \\ \text{CO}_2 \text{H} \end{array}$	(q)	Positional	

(c)	NO ₂ , NO ₂	(r)	Functional
(d)	OH CH ₂ OH	(s)	Metamers

43. Find sum of stereoisomer of following compound.

Number of stereoisomers

(a)
$$HO$$
 CO_2H
 CO_2H
 CO_2H
 CO_2H
 CH_3
 CO_2H
 OH
 OH

44.

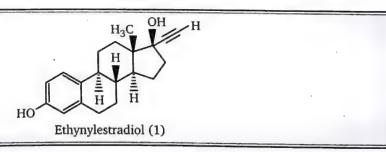
Prostaglandin E_1 1 is a compound produced by the body to regulate a variety of processes including blood clotting, fever, pain and inflammation.

- **A.** Which of the following functional groups is not contained in 1?
 - (a) A ketone
- (b) An alcohol
- (c) A carboxylic acid (d) An alkene

- (e) A nitrile
- B. How many asymmetric (stereogenic) centres are present in compound 1?
 - (a) 3
- (b) 4
- (c) 5
- (d) 6
- **C.** How many sp^2 hybridised carbon atoms are present in compound 1?
 - (a)
- (b) 2
- (c) 3
- (d) 4
- **D.** What is the geometric configuration about the double bond in compound 1?
 - (a) E

(b) Z

45.

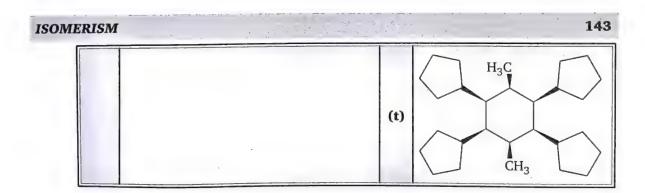


The synthetic steroid ethynylestradiol (1) is a compound used in the birth control pill.

- **A.** How many sp^3 hybridised carbon atoms are present in compound (1)?
 - (2)
- (b) 9
- (c) 10
- (d) 11 (e) 12
- **B.** How many sp^2 hybridised carbon atoms are present in compound (1)?
 - (-)
- (b) 5
- (c) 6
- (d) 7 (e) 8
- **C.** How many sp hybridised carbon atoms are present in compound (1)?
 - (a) 2
- (b) 4
- (c) 6
- (d) 8 (e) 10
- **D.** Which of the following functional group is contained in compound (1)?
 - (a) A ketone
- (b) An alcohol
- (c) A carboxylic acid (d) An ester
- E. How many asymmetric (stereogenic) centres are present in compound (1)?
 - (a) 2
- (b) 3
- (c) 4
- (d) 5

46. Match the column.

Water	the column.		A
(a)	Column (1) C ₂ -axis of symmetry	(p)	Column (II)
(p)	C ₃ -axis of symmetry	(p)	
(c)	Plane of symmetry	(r)	H ₃ C,
(d)	Centre of symmetry	(s)	



SUBJECTIVE PROBLEMS

1. Number of chiral isomers are:

Number of stereoisomer are

3. (i)
$$CH_3$$
 (ii) $H_2 \rightarrow (D)$ (Major). (C) Sum of number of stereoisomer (C) Degree of unsaturations in (D).

- 4. How many 5 membered parent chain alkane are possible for C₇H₁₆?
- 5. Theoretical possible geometrical isomer of

- **6.** Total number of possible structural isomers of C₅H₁₁Br.
- 7. Total number of plane of symmetry present in given compound is

- 8. Total number of isomers for C₄H₆Br₂ containing cyclobutane ring are (including stereoisomer)?
- 9. Total number of structural isomers of C₉H₁₈ containing cyclohexane ring.
- 10. How many structural isomer are possible for $C_4H_{10}O$ (only alcohol).
- 11. Number of structural isomer of C₆H₁₄ is .

12. (a)
$$CH_2$$
—Cl CH_2 —Cl (Number of plane of symmetry) CH_2 —Cl

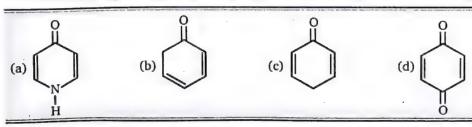
(b)
$$Cl \longrightarrow (y)$$
 (Number of mesoisomer of 1, 2-dichlorocyclopentane)

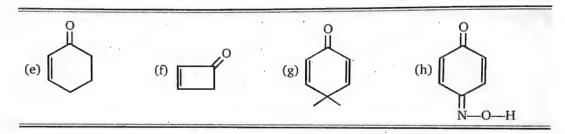
Sum of
$$(x+y=?)$$

13. Find out the total number of stereocentre in the given compound.

14. Find out the total number of stereoisomers of the given following compound.

- 15. Find the total number of isomers of C₇H₁₄ (only 5-membered ring).
- **16.** x = number of compounds which undergoes Tautomerisation to form an Aromatic product.





17. If molecule is pyramidal, X stereoisomers are possible for :

Cabed

find the value of **X**.

ANSWERS — LEVEL 2

2. a-s; b-r; c-q; d-p

10. a-p; b-q; c-s; d-r

14. a-s; b-r; c-q; d-p

12. a-q, r; b-p; c-p; d-q, r

4. a-p, q; b-p, q; c-p, q; d-p, r

6. a - p, r; b - q, s; c - q, r; d - p, s

8. a-r, s; b-p, q; c-r, s; d-p, r, s

1.
$$a-q$$
; $b-p$; $c-r$; $d-s$

3.
$$a-p$$
, s; $b-q$, r, s; $c-q$, r, s; $d-p$, s

5.
$$a-r$$
; $b-r$; $c-p$; $d-s$

7.
$$a-q$$
, r; $b-q$, s; $c-p$, q, r; $d-q$, s

9.
$$a-q$$
; $b-q$, s; $c-p$, q, s; $d-q$, s

11.
$$a-p$$
; $b-q$; $c-r$; $d-s$

13.
$$a-r$$
; $b-s$; $c-r$; $d-p$

15.
$$a-q$$
, r ; $b-r$, s ; $c-q$, r ; $d-p$, q

16.
$$(a-p-x)$$
; $(b-q, r-y)$; $(c-p-x)$; $(d-q, r-w)$

17.
$$A - b,h$$
; $B - a, g$; $C - c, e$; $D - d, f$

18.
$$A - d; B - a$$

19.
$$a+b+c+d=13$$

20.
$$A - b$$
, d , e ; $B - a$, c , f , h ; $C - i$, k , p

21.
$$A - e$$
, f, j; $B - a$, c, d, g, h, i, b; $C - None$

22.
$$w + x + y + z = 12$$

27.
$$A-d, h; B-d; C-f, h; D-h$$

28.	Compound	Α	В	С
	I	c	a	a
	II	С.	b	ъ
	III	c	a	a
	IV	С	b	b
	V	c	a	a
	VI	С	a	a
	VII	c	a	a·
	VIII	e	Ь	b
	IX	e	a	a
	X	e	b	b
29.	(a) (R)	(b) (L	

30A. 1 - N; 2 - Z; 3 - E; 4 - Z; 5 - Z; 6 - E; 7 - N

B. (a) 1 - Z; 2 - E; 3 - E; 4 - Z; 5 - Z; 6 - E

(b) 2⁹

31. 1-Z; 2-N; 3-E

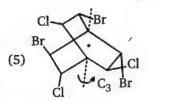
32. (a) A - 10, B - 4, C - 2

(c) A-4, B-2, C-0

(e) A-5, B-2, C-1

(g) A-4, B-2, C-0Me_ _Cl ...C₂

Me **33.** (1) Me



(7) C₃ axis, C₂ axis

(9) (C_3)

(12) C2-axis

(b) A - 5, B - 0, C - 5

(d) A - 2, B - 1, C - 0

(f) A-4, B-1, C-2

(h) A-4, B-1, C-2

$$(2) \underbrace{Et}_{Et} \underbrace{C_3^{Et}}_{C_3}$$

(4)
$$Cl$$
 Cl Cl

(6)
$$Me_3C$$
 N CMe_3 CMe_3

(8) C₃-axis

(13) C₂-axis

35.

Identical Enantiomer Diastereomer Constitutional Isomer

(4)
$$H \longrightarrow OH HO \longrightarrow H$$
 $H \longrightarrow CH_3$ \times \times \times

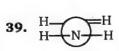
(5)
$$\stackrel{\text{H}}{\underset{\text{CH}_3}{\bigvee}} \stackrel{\text{CH}_3}{\underset{\text{H}}{\bigvee}} \stackrel{\text{CH}_3}{\underset{\text{CH}_3}{\bigvee}} \times \times \times \times \times \times$$

(9)
$$\times$$
 \times \times \times

(10)
$$\times$$
 \times \times \times \times \times

37.
$$a-p$$
; $b-q$; $c-r$; $d-s$

38.
$$a-s$$
; $b-p$; $c-q$; $d-r$





(Resonance)

 π -(vacant-p-orbital)

40. (1) 16

(2) 4

(3) 16

(4) 4

(5) 4

(6) 4

(7) 4

(8) 3

(9) 4

(10) 4

(11) 2

41. a-2, b-4, c-3, d-4, e-8, f-2

42. a-s; b-p; c-q; d-r

43. $a-2^5$, $b-2^5$, $c-2^7+2^3$, $d-2^9$

44. A - e; B - b; C - d; D - a

45. A - e; B - c; C - a; D - b; E - d

46. a-p, q, s, t; b-p, r; c-p, q, r, s, t; d-p

Subjective Problems

1. 3 (c, f, h)

2. 64

3. 5

4.5

5. 2

6. 8.

7. 3

151

8. 6

9. 12

10. 4

11. 5

12. 4

13. 4

14. 4

15. 8

16. 4 (a, b, c, h)

17. 6

GRIGNARD REAGENT



LEVEL-]

1. What is the major product of the following reaction?

$$\begin{array}{c}
CH_3MgI \\
Et_2O
\end{array}
\xrightarrow{H_3O^{\oplus}}$$

(a)
$$CH_3$$
 $CH_2 - OH$

(b)
$$CH_2 - CH_3$$

(c)
$$CH_3$$

2.
$$C - OCH_3$$
 $O \xrightarrow{PhMgBr} (P)$. Product P is:
$$CH_2 - COCH_3$$
 O

$$(a) \qquad \begin{array}{c} CH_3 O \\ C-Ph \\ CH_2 - COCH_3 \end{array} \\ (b) \qquad \begin{array}{c} CH_3 O \\ C-Ph \\ CH_2 - COCH_3 \end{array} \\ (c) \qquad \begin{array}{c} CH_3 C \\ C-Ph \\ CH_2 - COCH_3 \end{array} \\ (d) \qquad \begin{array}{c} CH_3 C \\ C-CH_3 C \\ C-CH_2 - C-CH_3 \end{array} \\ (d) \qquad \begin{array}{c} CH_3 C \\ C-CH_3 C \\ C-CH_2 - C-Ph \end{array} \\ (d) \qquad \begin{array}{c} CH_3 C \\ C-CH_3 C \\ C-CH_3$$

What is the ratio of (x/y) in above problem?

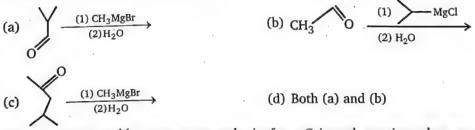
(a) 1.5

(b) 2

(c) 2.5

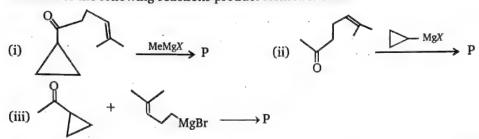
(d) 3

4. In which of the following reaction 2° alcohol is obtained as a product?



5. What product would you expect to obtain from Grignard reaction when an excess of phenylmagnesium bromide reacts with dimethyl carbonate CH₃OCOOCH₃ ?

6. In which of the following reactions product formed is same?



- (a) (i) and (ii)
- (b) (ii) and (iii)
- (c) (i) and (iii)
- (d) (i), (ii) and (iii)
- 7. Which of the following reaction sequences would be the best for synthesis of 2-pentanone?

(a)
$$CH_3 - CH_2 - CH_2 - C - H \xrightarrow{CH_3Mgl} \xrightarrow{H_3O^{\oplus}}$$

(b)
$$CH_2-CH_2 \xrightarrow{CH_3MgI} \xrightarrow{H_3O^{\oplus}} CH_3-CH-O$$

(c)
$$CH_3 - CH_2 - CH_2 - C \equiv N \xrightarrow{CH_3MgI} \xrightarrow{H_3O^{\oplus}}$$

(d)
$$CH_3 - CH_2 - C - H \xrightarrow{CH_3MgI} \xrightarrow{H_3O^{\oplus}}$$

8.
$$CO_2CH_3$$
 CO_2CH_3
 CO_2CH_3
 CCO_2CH_3
 $CCO_$

Number of moles (x) of Grignard reagent consumed in the above reaction is:

- (a) 2
- (b) 3
- (c) 4
- (d) 5

Which of the following combinations can not be used to prepare alcohol given above?

- (a) PhMgBr + 2-butanone $\xrightarrow{\text{NH}_4\text{Cl}}$
- (b) EtMgBr + Ph C CH₃ $\xrightarrow{\text{NH}_4\text{Cl}}$
- (c) $CH_3MgBr + Ph C Et \xrightarrow{NH_4Cl}$
- (d) EtMgBr + Ph C -CH₂ -CH₃ $\xrightarrow{\text{NH}_4Cl}$

10. Et—O—C—O— Et—
$$(1) CH_3MgBr (excess)$$
 (A), Product (A) is:

(d)
$$CH_3 - CH_2 - CH_3$$

11. PhMgBr +
$$O \longrightarrow O \longrightarrow O \longrightarrow (A)$$
. Product (A) is:

12.
$$\bigcirc \bigcap_{C - NOCH_3}^{O} + CH_3 MgBr \xrightarrow{5\% HCl}_{\text{Methyl magnesium bromide 75 equivalent}} (P) \cdot \text{Product } (P) \text{ is :}$$

N-Methoxy-N-methyl benzamide 1-equivalent

13. Point out the incorrect synthesis:

$$\begin{array}{c}
0\\ \parallel\\ (1) \text{ CH}_3-\text{C-CH}_3
\end{array}$$

$$\begin{array}{c}
(2)\text{H}^+
\end{array}$$

(c)
$$PhMgBr + \xrightarrow{(1)} OH \xrightarrow{(2)} Ph - CH_2 - CH_2 - OH$$

(d)
$$PhMgBr + \frac{(1) HCHO}{(2) H^+} Ph - CH_2 - OH$$

14.
$$CH_3 - C - O - H \xrightarrow{NaHCO_3} (A) \xrightarrow{(i) PhMgBr} (B)$$

$$CH_{3} - S - O - H \xrightarrow{\text{NaH CO}_{3}} (C) \xrightarrow{\text{(i) PhMgBr}} (D)$$

Product (B) and (D) in the above reaction are:

(a)
$$Ph - C - O - H$$
, $Ph - S - O - H$ (b) $Ph - C - O - H$, $Ph - S - O - H$

15. OH OEt
$$(2)$$
 (A) ; Product (A) in this sequence is :

16.
$$CH_3 MgBr \rightarrow NH_4Cl \rightarrow Products.$$

Comment on optical activity of the products. They are ;

- (a) racemic mixture
- (b) diastereomers

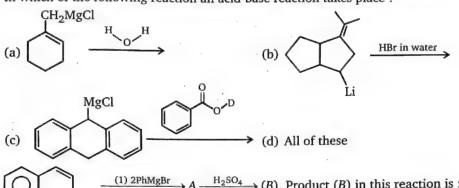
(c) meso forms

(d) optically inactive due to absence of chiral centre

Grignard Reagent

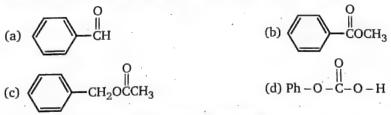
157

In which of the following reaction an acid-base reaction takes place?



(B), Product (B) in this reaction is: (2) H₂O

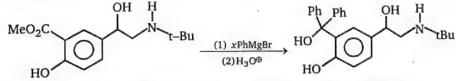
All of the following compounds react with ethylmagnesium bromide. Alcohols are formed from three of the compounds. Which one does not give an alcohol?



- A student was carrying out a Grignard reaction between PhMgBr and ethyl benzoate. She ran 20. out of anhydrous ether just after the Grignard reagent was made. Which of the following solvents can still be used to dissolve the ethyl benzoate for its reaction with already formed PhMgBr?
 - (a) acetone

21.

- (b) ethyl acetate
- (c) absolute alcohol (d) benzene



Number of equivalents of Grignard reagent (x) used in reaction (1) is:

- (a) 3 equivalent
- (b) 4 equivalent
- (c) 5 equivalent
- (d) 6 equivalent

24.

22.
$$\begin{array}{c}
\text{OH} \\
\text{CH - Ph} \\
\text{CH - Ph} \\
\text{CH - Ph} \\
\text{CH - Ph} \\
\text{CH - Ph}
\end{array}$$

The given product can not be obtained in the above reaction. Identify the correct product obtained.

(a)
$$\bigcap_{N \to CH - Ph} (b) \bigcap_{N \to CH - Ph} (c) \bigcap_{N \to CH - Ph} (d) \bigcap_{N \to CH - Ph} (d)$$

23. Which of the following gives two isomers of 3° alcohol, when treated with phenyl magnesium bromide?

(a) (b) (c) (d)
$$\stackrel{D}{\longrightarrow}$$
 Me $\stackrel{MeMgBr}{\longrightarrow}$ Product of the reaction is :

(a) Me H (b) Me H (c) Both (a) and (b) (d) None of these

25.
$$\xrightarrow{RMgBr}$$
 Product; The product of the reaction is:

(a) $HSO_2 - CH_2 - CH_2 - CH_2 - CH_2 - R$ (b) $H - SO_2(CH_2)_3 - R$

(c) $MgBr$ (d) $H - SO_2(CH)_3 - R$

26. When carboxylic acid reacts with organolithium reagents to give ketones, side reaction sometimes occur. For example,

Value of (x) in above reaction is:

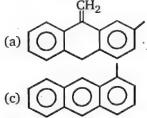
- (a) 2
- (b) 3
- (c) 4
- (d) 5
- **27.** Which of the following alcohol can not be prepared by the reaction of acid chloride with excess of Grignard reagent followed by acidification?

(a)
$$Ph \xrightarrow{OH} Ph$$
 CH_3

- (b) Ph CH
- (c) Et Et
- (d) CH₃ Et

28.
$$(i) \text{ MeMgI} \atop (ii) \text{ NH}_4\text{Cl}} (A) \xrightarrow{\text{H}^+} (B) \atop (\text{major})$$

Product (B) of the above reaction is:



- 29. The reaction of elemental sulphur with Grignard reagent followed by acidification leads to the formation of
 - (a) mercaptan
- (b) sulphoxide
- (c) thioether
- (d) sulphonic acid

+ MgBrCH₂CH₂CH₂CH₂MgBr $\xrightarrow{(i)THF}$ product; Product of the reaction is :

(a)
$$HO = (CH_2)_3 = C = CH_2 = CH_2 = CH_3$$
 (b)

31.
$$\bigcirc C - Cl \xrightarrow{(i) \times CH_3MgBr} \bigcirc C - CH_3$$

$$C - Cl \xrightarrow{(ii) H^+} \bigcirc CH_3$$

$$Me \xrightarrow{C} OH$$

$$Me$$

$$Me$$

Number of moles of CH₃MgBr consumed in above reaction is :

(a) 2

(b) 4

(c) 6

(b)

. (d) 8

32. End product of the given reaction is:

(a)
$$CI \xrightarrow{Mg} (A) \xrightarrow{1.CH_2O} (B)$$
(b) CH
(c) CH
(d) CH

33. Which of the following compound is not a suitable solvent for Grignard reaction?

(a)
$$\bigcup_{O \text{ (THF)}}$$

(c) $CH_3 - O - CH_2 - CH_2 - O - CH_3$

34. $\underset{\text{(2 mole)}}{\text{Br}} \xrightarrow{\text{Mg}} \text{Predict major product of the reaction}$:

(a) /

(b) /

(c) //

(1, 4-dioxane)

(d) /

35. Which of the following reaction sequences would be the best for synthesis of t-butyl alcohol?

(a)
$$CH_3CH_2MgBr + CH_2 - CH_2 \xrightarrow{Et_2O} \xrightarrow{H_3O\oplus}$$

(b)
$$CH_3CH_2CH_2MgBr \xrightarrow{CO_2} \xrightarrow{H_3O\oplus}$$

(c)
$$CH_3MgBr + CH_3 - C - CH_3 \xrightarrow{Et_2O} \xrightarrow{H_3O^{\oplus}}$$

(d)
$$CH_3CH_2MgBr + CH_3 - C - H \xrightarrow{Et_2O} \xrightarrow{H_3O^{\oplus}}$$

What is the major product of the following reaction?

$$CH_3 - C \equiv N \xrightarrow{CH_3MgI} \xrightarrow{H_3O^{\oplus}}$$

(a)
$$CH_3 - CH_2 - NH - CH_3$$

(c) $CH_3 - C - CH_3$

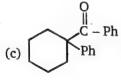
- $\xrightarrow{\text{(1) PhMgBr}} \text{Products; Product obtained in this reaction are :}$
 - (a) diastereomers
- (b) racemic
- (c) pure enantiomer (d) meso
- **38.** $CH_3CO_2Et + (CH_2)_5(MgBr)_2 \xrightarrow{(2) H^+} C_7 \underset{(A)}{H_{14}}O$; compound (A) will be:

(c)
$$CH_3 - C - (CH_2)_4 - CH_3$$

-OH

39.
$$(1) \xrightarrow{\text{PhMgBr/CuCl}} (A) ; A \text{ will be} :$$

(a)
$$Ph$$
 (b) $C-I$



$$(d) \bigcap_{Ph} C - Ph$$

40. MeO OMe
$$\frac{n-Bu_2Cu_{Li}}{(n-Bu=n-butyl \text{ group})}$$
 Product of the reaction will be:

162

ORGANIC Chemistry for IIT-JEE

(?); Product of this reaction is:



42. Ethyl acetoacetate when reacts with one mole methyl magnesium iodide then product of reaction will be:

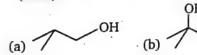
(a)
$$CH_3 - C - CH_2 - C - CH_3$$

(c)
$$CH_3 - C - CH - CO_2Et$$

$$MgBr$$

(d)
$$CH_2^- - C - CH_2 - CO_2Et$$

43. $CH_3MgBr + Et - O - C - O - Et \xrightarrow{(2) H^{\oplus}} (A)$; Product A is:







- For the sequence of reactions, $A \xrightarrow{C_2H_5MgI} B \xrightarrow{H_2O/H^+} B$ tert-Pentyl alcohol. The compound A in the sequence is:
 - (a) 2-Butanone
- (b) Acetaldehyde (c) Acetone
- (d) Propanal
- **45.** PhMgBr + CH₃ CN $\xrightarrow{\text{H}_3O^{\oplus}}$ (A) Ph C O H $\xrightarrow{\text{(1) excess CH}_3-\text{Li}}$ (A)

Same product (A) will form in both reactions. A is:

(a) Ph
$$-C - CH_3$$

Same product (A) will form in both reactions. A is : OH OH OHO (c) Ph — C — CH
$$_3$$
 (d) Ph — CH $_2$ — CO $_2$ H CH $_3$

- Which of the following Grignard reagent can be prepared?
 - (a) $Br Mg CH_2 CH_2 CH_2 O H$ (b) $Br Mg CH_2 CH_2 SH$
 - (c) $BrMg CH_2 CH_2 NH_2$
- (d) $BrMg CH_2 CH_2 N CH_3$
- In the reaction sequence: 47.

$$(i) CH_3MgBr/CuCl / (ii) H_2O/H^+ (X), Product (X) will be :$$

- $CH_3MgBr(excess)$ $\rightarrow A$. A (alcohol) can also be obtained by : $(C_2H_5O)_2CO$ H₃O⁺
 - (a) CH₃CH₂CHO-

- (d) as in (b) and (c)
- The principal product of the reaction between methyl butanoate and 2 moles of CH_3MgBr 49. after hydrolysis is:
 - (a) C₃H₇COCH₃

(b) $C_3H_7C(OH)(CH_3)_2$

(c) C₃H₇CHOHCH₃

- (d) $C_3H_7COCH(CH_3)_2$
- Which of the following compounds will form hydrocarbon on reaction with Grignard reagent? 50.
 - (a) CH₃CH₂OH
- (b) CH₃CHO
- (c) CH₃COCH₃
- (d) CH₃CO₂CH₃
- What is the product (B) of the following reaction sequence?

- **52.** Which, if any, of the following pairs of reagents could be used to prepare 2-phenyl-2-butanol?
 - (a) $CH_3CH_2MgBr + Ph C CH_3$
- (b) CH₃CH₂MgBr + C₆H₅CH₂CH
- (c) $CH_3MgI + C_6H_5CH_2CCH_3$
- (d) $C_6H_5MgCl + CH_3CCH_2CH_2CH_3$
- **53.** What is the product of the following reaction?

$$O + 2CH_3MgBr \xrightarrow{1. \text{ diethyl ether}} Product$$

- (a) HO CHC H₂CH₂CH₂CH OH
- (b) $\mathrm{CH_3OCH_2CH_2CH_2CH_2CHCH_3}$
- (c) HOCH₂CH₂CH₂CH₂CH₂C OH
- (d) $HOCH_2CH_2CH_2CH_2CHOCH_3$ CH_3
- 54. $\xrightarrow{\text{Ph}} \xrightarrow{\text{Mg}} A \xrightarrow{\text{1. HCHO}} B$; Product (B) is :

(a)
$$Ph$$
 CH_2-OF

- (b) $Ph C \equiv C CH_2 CH_2 CH_2 OH$
- (c) $Ph C \equiv C CH_2 CH_2 OH$
- (d) $Ph CH_2 C \equiv C CH_2 CH_2 OH$
- **55.** What sequence of steps represents the best synthesis of 4-heptanol (CH₃CH₂CH₂)₂CHOH?
 - (a) $CH_3CH_2CH_2MgBr(2moles) + formaldehyde(H_2C = O)$ in diethyl ether followed by $H_3O + CH_2CH_2MgBr(2moles) + formaldehyde(H_2C = O)$
 - (b) $CH_3CH_2CH_2MgBr + butanol (CH_3CH_2CH_2CH = O)$ in diethyl ether followed by H_3O^+
 - (c) $CH_3CH_2CH_2CH_2MgBr + acetone[(CH_3)_2C = O]$ in diethyl ether followed by H_3O^+
 - (d) $(CH_3CH_2CH_2)_2CHMgBr + formaldehyde (H_2C = O)$ in diethyl ether followed by H_3O^+
- 56. Et $(1) \text{ MeMgBr} \longrightarrow \text{Comment on stereochemistry of products}$:
 - (a) diastereomers

(b) racemic

(c) single stereoisomer

(d) meso

57.
$$CH \longrightarrow CH_3MgBr \longrightarrow xCH_4$$
 $CH_2 \longrightarrow SH$
 $(Excess)$

What is the value of x in the above reaction?

(a) 1

(b) 2

(c) 3

- (d) 4
- 0.40 g of an organic compound (A), (M.F.- C_5H_8O) reacts with x mole of CH_3MgBr to 58. liberate 224 mL of a gas at STP. With excess of H2, (A) gives pentan-1-ol. The correct structure of (A) is:

(a)
$$CH_3 - C \equiv C - CH_2 - CH_2 - OH_2$$

(b)
$$CH_3 - CH_2 - C \equiv C - CH_2 - OH$$

(c)
$$H - C \equiv C - CH_2 - CH_2 - CH_2 - OH$$

(d)
$$H - C \equiv C - CH_2 - CH_3$$

59.
$$CH_3 - CH = CH_2 \xrightarrow{Br_2} \xrightarrow{Mg} \xrightarrow{CH_3 - C - CH_3} \xrightarrow{H^+} \xrightarrow{\Delta} (X)$$
(low conc.) (major)

End product (X) of the above reaction is:

$$\begin{array}{c} \operatorname{CH_2} \\ \parallel \\ \operatorname{CH}_2 = \operatorname{CH} - \operatorname{CH_2} - \operatorname{C} - \operatorname{CH_3} \end{array}$$

(b)
$$H_2C = CH - CH = C - CH_3$$

 CH_3

(c)
$$H_2C = CH - CH_2 - C - CH_3$$

 CH_3

(d)
$$H_2C = CH - CH_2 - CH - CH_2 - OH$$
 CH_3

60.
$$CH_2 - CH - CH_2 - Br \xrightarrow{Mg} (A) \xrightarrow{CH_3I} (B)$$
; Product (B) is:

(a)
$$CH_2 - CH - CH_2 - CH_3$$

(b)
$$CH_3 - O - CH_2 - CH_2 - CH_3$$

(c)
$$H_2C = CH - CH_2 - O - CH_3$$
 (d) $H_2C - CH - CH_3$

(d)
$$H_2C - CH - CH$$

61. Compound A was treated with a large excess of CH₃MgBr. The resulting product was exposed to POCl₃/pyridine to give compound B, as one of many products:

Which of the following compounds can be A?

62. Identify product Z in the following reaction sequence:

$$H_2C = CHCH_2Br \xrightarrow{NaCN} Y \xrightarrow{1. C_6H_5MgBr, diethylether} Z$$

(a)
$$H_2C = CHCH_2CC_6H_5$$

(b)
$$H_2C = CHCH_2NHCC_6H_5$$

(c)
$$H_2C = CHCH_2CHC_6H_5$$

(d)
$$H_2C = CHCH_2CHC_6H_5$$

(Consider all steps and intermediate) correct statement is:

(a) Nucleophilic addition

63.

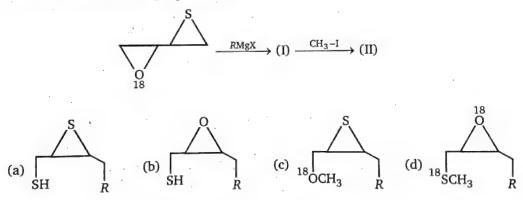
(b) Nucleophilic substitution reaction

(c) Product obtained is chiral

(d) All

64. Which combination(s) of alkyl bromide and epoxide can be used to prepare the following product by addition of the Grignard reagent derived from the alkyl bromide to the epoxide?

65. What will be the final major product?



66. Give the expected product of the following reaction.

	ANSWERS — LEVEL 1														
1.	(b)	2.	(d)	3.	(b)	4.	(d)	5.	(c)	6.	(d)	7.	(c)	8.	(c)
9.	(d)	10.	(c)	11.	(b)	12.	(b)	13.	(b)	14.	(c)	15.	(b)	16.	(b)
17.	(d)	18.	(d)	19.	(d)	20.	(d)	21.	(c)	22.	(b)	23.	(b)	24.	(c)
25.	(c)	26.	(b)	27.	(d)	28.	(d)	29.	(a)	30.	(b)	31.	(b)	32.	(b)
33.	(d)	34.	(b)	35.	(c)	36.	(c)	37.	(b)	38.	(b)	39.	(b)	40.	(a)
41.	(b)	42.	(c)	43.	(b)	44.	(c)	45.	(c)	46.	(d)	47.	(b)	48.	(d)
49.	(b)	50.	(a)	51.	(a)	52.	(a)	53.	(c)	54.	(b)	55.	(b)	56.	(a)
57.	(c)	58.	(c)	59.	(b)	60.	(c)	61.	(d)	62.	(a)	63.	(d)	64.	(b)
65.	(c)	66.	(d)				,								



1. Comprehension

Grignard reagent is usually prepared by

$$R - X + Mg \xrightarrow{\text{Et}_2O} RMgX$$

Grignard reagent

$$Ar - X + Mg \xrightarrow{Et_2O} ArMgX$$

Grignard reagent

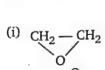
Grignard reagent acts as a strong base. Grignard reagent carry out nucleophilic attack in absence of acidic hydrogen. Grignard reagent form complex with its ether solvent. Complex formation with molecule of ether is an important factor in the formation and stability of Grignard reagent.

- A. What is the correct order of reactivity of halides with magnesium?
 - (a) R Cl > R Br > R I
- (b) R Br > R Cl > R I
- (c) R I > R Br > R Cl
- (d) R I = R Br = R Cl
- B. Which of the following will undergo acid-base reaction with Grignard reagent?
 - (a) $HC \equiv CH$

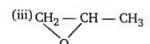
(b) R −OH

(c) $R - CO_2H$

- (d) All of these
- **C.** Which of the following reactants give primary alcohol as a major product when reacts with *RMgX* followed by acidification?



(ii) H-C-H



- (iv) CH₃ C H
- (v) O
- (vi)

- (a) i, ii, v
- (b) i, ii, v, vi
- (c) ii, iv, vi
- (d) v, iv, iii, vi
- **D.** Cl C O Et $\xrightarrow{\text{(1) } x \text{RMgX}}$ 3° alcohol. Value of x is :
 - (a) 2
- (b) 3
- (c) 4
- (d) 5
- E. $H-O-CH_2-CH_2-C-O-Et \xrightarrow{(1) \times PhMgBr} HO-CH_2-CH_2-C-Ph$, Value of x is:
 - (a) 2
- (b) 3
- (c) 4
- (d) 5

F. Which of the following Grignard reagents is not possible?

(a) $HS - CH_2 - CH_2 - CH_2MgBr$

(b) $HO - CH_2 - CH_2 - CH_2MgBr$

(c) $NH_2 - CH_2 - CH_2 - CH_2MgBr$

(d) All of these

G. How many different Grignard reagents when react with EtOH, give *n*-butane as product (excluding stereoisomerism).

(a) 1

(b) 2

(c) 3

(d) 4

2. Match the column I and II. (Matrix)

*	Column (I)		Column (II)		
	Reactant	Product			
(a)	$ \begin{array}{c} O \\ \\ PhMgBr + Cl - C - O - Et \longrightarrow \\ \text{(excess)} \end{array} $	(p)	Ph – CH ₂ – OH		
(b)	$ \begin{array}{c} O \\ \\ PhMgBr + H - C - O - Et \xrightarrow{H^{\oplus}} \end{array} $ (excess)	(q)	Ph - CH - Ph OH		
(c)	O PhMgBr+ H − C − H — H [⊕] → (excess)	(r)	OH 		
(d)	$ \begin{array}{c} O \\ \\ PhMgBr + CH_3 - C - O - Et \xrightarrow{H^{\oplus}} \end{array} $ (excess)	(s)	OH 		

3. Match the column I and II. (Matrix)

at some from	Column (I)	· san yan dan dan dan dan dan dan dan dan dan d	Column (II)
	Reaction		Reactant
(a)	PhMgBr + (A) \longrightarrow 1°alcohol	(p)	$\begin{array}{c c} O & O \\ \parallel & \parallel \\ CH_3 - C - CH_2 - C - CH_3 \end{array}$
(b)	PhMgBr + (B) \longrightarrow H^{\oplus} \rightarrow 2°alcohol	(q)	O CH ₃ - C - CH ₃
(c)	PhMgBr + (C) \longrightarrow 3°alcohol	(r)	O CH ₃ -C-H
(d)	$PhMgBr + (D) \xrightarrow{H^{\oplus}}$	(s)	O H-C-H

Match the missing reactant A, B, C, D

4. Match the column I and II. (Matrix)

سيعتبس	Column (I)	Col	Column (II)		
	Reaction	Moles of PhMgBr used			
(a)	$\begin{array}{c} O \\ \\ PhMgBr + Et - O - C - O - Et \xrightarrow{H^{\oplus}} 3^{\circ}alcohol \end{array}$	(p)	1		
(ь)	PhMgBr + HO - CH ₂ - C - CH ₃ $\xrightarrow{H^{\oplus}}$ 3° alcohol	(q)	2		
(c)	$\begin{array}{c} O \\ \\ PhMgBr + CH_3 - C - CH_3 \xrightarrow{H^{\oplus}} 3^{\circ} alcohol \end{array}$	(r)	3		
(d)	PhMgBr + $C - Cl$ HO HO HO HO $H\oplus$ HO HO $H\oplus$ HO $H\oplus$ HO HO $H\oplus$ HO HO $H\oplus$ HO	(s)	4		

5. When 20 g of a compound (A) (M.F. = $C_4H_{10}O_4$) reacts with excess of CH_3MgBr , 14.6 L of CH_4 is obtained at STP. What is structural formula of (A) ?

CH₂OH

OH

$$CH_2$$
—OH

(a) $(CH - OH)_2$ (b) $HO - CH_2$ —

 CH_2 —OH

(c) HO OH

(d) Both (a) & (b)

SUBJECTIVE PROBLEMS

1.

$$OH \xrightarrow{O} H \xrightarrow{excess} H^{\oplus} \xrightarrow{H^{\oplus}} \Lambda (X)$$

How many geometrical isomer of (X) is possible?

2. How many isomer of C_4H_8O when reacts with CH_3MgBr followed by acidification to give 2° alcohol (only consider carbonyl isomers)?

(including stereoisomer)

3.

Total number of RMgX are consumed in the following reaction

$$Cl = N$$

$$C = N$$

- **4.** How many isomers of $C_4H_{10}O$ reacts with CH_3MgBr to evolve CH_4 gas ? (Excluding stereoisomer)
- 5. How many carbonyl isomers of $C_5H_{10}O$ which reacts with PhMgBr to give racemic mixture?
- 6. How many moles of Grignard reagent will consume when it reacts with following compound?

ANSWERS — LEVEL 2

1.
$$A-c$$
; $B-d$; $C-a$; $D-b$; $E-b$; $F-d$; $G-b$

" # R79"

2.
$$a-r$$
; $b-q$; $c-p$; $d-s$

3.
$$a-s$$
; $b-r$; $c-q$; $d-p$

4.
$$a-r$$
; $b-q$; $c-p$; $d-s$

5. (d)

Subjective Problems

1. 4

2. 2

3. 7

4. 4

5. 5

6 Q



HYDROCARBONS (ALKANES)



- 1. On halogenation, an alkane gives only one monohalogenated product. The alkane may be:
 - (a) 2-methyl butane

(b) 2, 2-dimethyl propane

(c) cyclopentane

- (d) both (b) and (c)
- 2. Which of the following compounds can be best prepared by Wurtz-reaction?
 - (a) Iso-butane

(b) *n*-butane

(c) n-pentane

- (d) Iso-pentane
- **3.** A hydrocarbon A (V.D. = 36) forms only one monochloro substitution product. A will be:
 - (a) iso-pentane

(b) neo-pentane

(c) cyclohexane

- (d) methyl-cyclohexane
- **4.** Ethyl iodide and *n*-propyl iodide are allowed to undergo Wurtz reaction. The alkane which will not be obtained in this reaction is:
 - (a) butane

(b) propane

(c) pentane

(d) hexane

5.
$$CH_3$$
— CH — CH_2 — CH_3 — CH_3 — CH_2 — CH_3 —

Number of chiral centers generated during monochlorination in the above reaction :

- (a) 1
- (b) 2
- (c) 3
- (d) 4

 $CH_3Cl \longrightarrow CH_4$

Above conversion can be achieved by:

(a) Zn / H⁺

- (b) LiAlH₄
- (c) Mg / (ether) then H₂O
- (d) all of these

n-Butane $\frac{Cl_2/hv}{}$ 7.

> Give the total number of monochloro products (including stereoisomers), which are possible in the above reaction.

- (b) 3
- (c) 4
- (d) 5

 $CH_4 + Cl_2 \xrightarrow{hv} CH_3Cl + HCl$

To obtain high yields of CH₃Cl, the ratio of CH₄ to Cl₂ must be :

(a) high

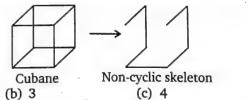
(b) low

(c) equal

- (d) can't be predicted
- Double bond equivalent of cubane is:

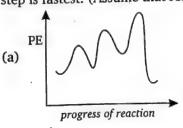


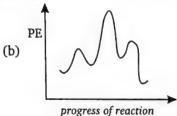
- (a) 4
- (b) 5
- (c) 6
- (d) 7
- How many bond cleavages are required to convert cubane into non-cyclic skeleton? 10.

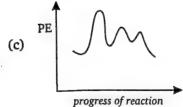


- (a) 2

- (d) 5
- Draw an energy profile diagram for a three step reaction in which first step is slowest and last 11. step is fastest. (Assume that reaction is exothermic)







(d) None of these

(d) 7

- 12. CH_3 —CH— CH_2 — CH_3 $\xrightarrow{Cl_2}$ \xrightarrow{hv} (x) = Number of monochloro product including CH_3 stereoisomers.
- (a) 4 (b) 5 $\xrightarrow{\text{ND}_2 \text{ND}_2} (P)$ 13.

Product (P) is:



- (p) D
- (c) D

(c) 6

- (d) both (b) & (c)
- Double bond equivalent (degree of Unsaturation) of (A) is:

 (a) 1 (b) 2 (c) 3 (d) 4
- 15. Arrange the following alkanes in decreasing order of their heats of combustion.
 - (i) CH₃—C—CH₃
 CH₃
 (Neo-pentane) (i)

- (ii) CH₃— CH CH₂ CH₃

 CH₃

 (Iso-pentane) (ii)
- (iii) $CH_3 CH_2 CH_2 CH_3 CH_3$
- (a) (i) > (ii) > (iii)

(b) (iii) > (i) > (ii)

(c) (iii) > (ii) > (i)

- (d) (i) > (ii) > (ii)
- 16. $\begin{array}{c} H \\ CH_3 \end{array} + H_2 \xrightarrow{Pd}$

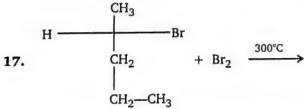
Product of the above reaction will be:

(a) Racemic mixture

(b) Diastereomers

(c) Meso

(d) Constitutional isomers



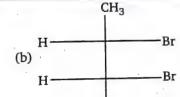
Which of the following compound will not be obtained as a product in the above reaction?

CH₃

H

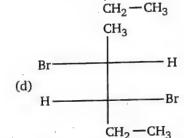
Br

H



 $H \xrightarrow{CH_3} Br$ (c) $CH_2 Br$

CH2-CH3



18. Following are the structures of four isomer of hexane. Among the names given below, which correctly identifies the fifth isomer?

CH₃CH₂CH₂CH₂CH₂CH₃ (CH₃)₂CHCH₂CH₂CH₃

 CH_3

 $\begin{array}{c} (\mathrm{CH_3})_3\mathrm{CCH_2CH_3} \\ (\mathrm{CH_3})_2\mathrm{CHCH(CH_3)_2} \end{array}$

- (a) 2-Methyl pentane
- (c) 2,3-Dimethyl butane
- (b) 2-Ethyl butane
- (d) 3-Methyl pentane
- 19. Which of the following describes the best relationship between the methyl groups in the chair conformation of the substance shown below?

(a) Trans

(b) Anti

(c) Gauche

- (d) Eclipsed
- **20.** Compare the stabilities of the following two compounds (A) and (B):

A: cis-1-ethyl-3-methyl cyclohexane

B: trans-1-ethyl-3-methyl cyclohexane

(a) A is more stable

- (b) B is more stable
- (c) A and B are of equal stability
- (d) No comparison can be made
- 21. Which conformation of ethane has the lowest potential energy?
 - (a) Eclipsed

(b) Skew

(c) Staggered

- (d) All will have equal potential energy
- **22.** Ethane is subjected to combustion process. During the combustion the hybrid state of carbon changes from :
 - (a) sp^2 to sp^3

(b) sp^3 to sp

(c) $sp to sp^3$

(d) sp^2 to sp^2

23.
$$CH_3 - CH_2 - CH_2 - CH_3 \xrightarrow{AlCl_3} CH_3 - CH - CH_2$$

Above reaction is an example of :

(a) isomerization

(b) polymerization

(c) cracking

- (d) de-hydrogenation
- 24. Which of the following has highest chlorine content?
 - (a) Pyrene
- (b) DDT
- (c) Chloral
- (d) Gammaxene

- 25. Pure methane can be prepared by:
 - (a) Wurtz reaction

- (b) Kolbe electrolysis method
- (c) soda-lime de-carboxylation
- (d) reduction with H₂
- 26. Calcium carbide + heavy water ---- ?

The product of the above reaction is:

- (a) C_2H_2
- (b) CaD₂
- (c) Ca(OD)₂
- (d) CD₄

27.

Ethyl cyclopentane

Ethyl cyclohexane

Ethyl cycloheptane

Arrange the compounds I, II and III in decreasing order of their heats of combustion:

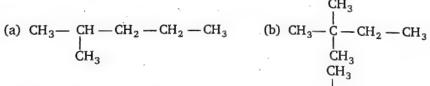
(a) II > I > III

(b) I > II > III

(c) III > II > I

- (d) III > I > II
- An alkane (mol. wt. = 86) on bromination gives only two monobromo derivatives (excluding stereoisomers). The alkane is:

(a)
$$CH_3$$
— CH — CH_2 — CH_3 — CH_3
 CH_3



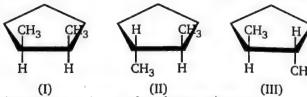
- 29. Order of the bond strength of C H bonds involving sp, sp^2 and sp^3 hybridized carbon
 - (a) $sp > sp^2 > sp^3$

(b) $sp^3 > sp^2 > sp$

(c) $sp^2 > sp^3 > sp$

(d) $sp^2 > sp > sp^3$

30.



Among the structures given, select the enantiomers:

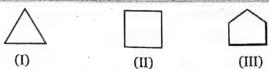
(a) I and II

(b) I and III

(c) II and III

(d) I, II and III

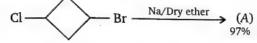
31.



The correct order of reactivity of I, II & III towards addition reactions is:

- (a) I > III > II
- (b) I > II > III
- (c) III > II > I
- (d) III > I > II

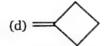
32.



Product (A) of above reaction is:



- (b)
- (c)

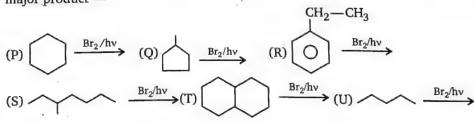


- **33.** Which of the following reactants is suitable for preparation of methane and ethane by using one step only?
 - (a) $H_2C = CH_2$

(b) CH₃OH

(c) CH₃ --- Br

- (d) $CH_3 CH_2 OH$
- **34.** How many carbon atoms does an alkane (not a cycloalkane) need before it is capable to exist in enantiomeric form?
 - (a) 4
- (b) 5
- (c) 6
- (d) 7
- **35.** Among the following free radical bromination reactions, select those in which 2° halide is the major product —



- (a) P, Q, R, S
- (b) P, R, U
- (c) P, R, S, T
- (d) P, Q, R, S, T

36. (A) + $Cl_2 \xrightarrow{hv}$ monochloro product

To maximise the yield of monochloro product in the above reaction?

- (a) Cl₂ must be added in excess
- (b) Reactant (A) must be added in excess
- (c) Reaction must be carried out in dark
- (d) Reaction must be carried out with equimolar mixture of Cl2 and A
- 37. $CH_3 CH_2 CH_2 CH_3 \xrightarrow{Br_2/h\nu}$

Major product in the above reaction is:

(a) Racemic mixture

(b) Meso

(c) Diastereomers

(d) Constitutional isomers

- 38. Select the chain propagation steps in the free-radical chlorination of methane.
 - (1) $Cl_2 \longrightarrow 2Cl^{\bullet}$

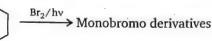
- (2) $Cl^{\bullet} + CH_4 \longrightarrow CH_3Cl + H^{\bullet}$
- (3) $Cl^{\bullet} + CH_4 \longrightarrow CH_3^{\bullet} + HCl$
- $(4) H^{\bullet} + Cl_2 \longrightarrow HCl + Cl^{\bullet}$
- (5) $CH_3^{\bullet} + Cl_2 \longrightarrow CH_3Cl + Cl^{\bullet}$
- (a) 2, 3, 5

(b) 1, 3, 6

(c) 3, 5

(d) 2, 3, 4

39. CH₃



The number of possible monobromo products is (excluding stereoisomers):

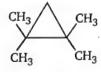
- (a) 4
- (b) 5
- (c) 8
- (d) 10

40.

$$H^{d}$$
 CH_{2}
 H^{b}
 CH_{2}
 H^{c}
 H^{b}
 H^{c}
 H^{c}

Br* will abstract which of the hydrogen most readily?

- (a) a
- (b) b
- (c) c
- (d) d
- 41. Arrange the following compounds in decreasing order of their heats of combustion :





(ii)



(a) (iii) > (ii) > (i)

(iii) (b) (ii) > (i) > (iii)

(c) (iii) > (i) > (ii)

- (d) (i) > (ii) > (iii)
- **42.** $CH_3 CH_2 CH_2 CH_2 F$

Arrange the hydrogens a, b, c, d, in decreasing order of their reactivities towards chlorination:

(a) a > b > c > d

(b) b > c > d > a

(c) b > c > a > d

- (d) c > b > a > d
- **43.** On catalytic reduction (H_2/Pt) how many alkenes will give *n*-butane?
 - (a) 1

(b) 2

(c) 3

- (d) 4
- **44.** On catalytic reduction (H₂/Pt) how many alkenes will give 2-methylbutane?
 - (a) 1

(b) 2

(c) 3

(d) 4

Cl₂ (excess)/hv 45.

How many dichloro products are formed in the above reaction (including stereoisomers)?

(a) 5

(b) 6

(c) 7

(d) 9

46.
$$CH_3 - CH$$
 $C = C < CH_3 \longrightarrow H_2/Pt$

Product of the above reaction will be:

- (a) Racemic mixture

(b) Diastereomers

(c) Meso

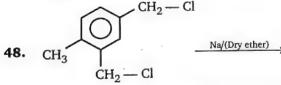
(d) Constitutional isomers

47.
$$Ph - CH_2 - CH - CH_3 \xrightarrow{Br_2/hv}$$

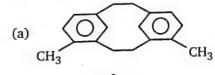
Product of the above reaction will be:

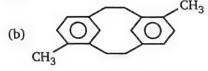
- (a) Diastereomers
- (c) Meso

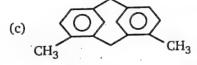
- (b) Racemic mixture
 - (d) Constitutional isomers



Products obtained in above Wurtz reaction is:







- (d) Both (a) and (b)
- Rank the transition states that occur during the following reaction steps in order of 49. increasing stability (least \rightarrow most stable):

1.
$$H_3C - OH_2 \longrightarrow CH_3^+ + H_2O$$

2.
$$(CH_3)_3C - \overset{+}{O}H_2 \longrightarrow (CH_3)_3C^+ + H_2O$$

3.
$$(CH_3)_2CH \stackrel{\circ}{-O}H_2 \longrightarrow (CH_3)_2CH^+ + H_2O$$

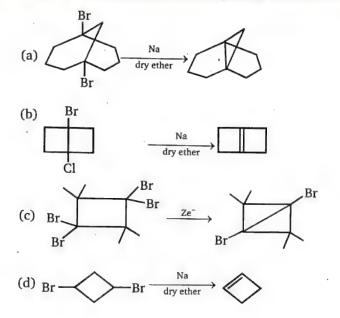
(a) 1 < 2 < 3

(b) 2 < 3 < 1

(c) 1 < 3 < 2

(d) 2 < 1 < 3

50. Which of the following does not represent major product of that reaction?



						ANSV	VERS	— LE	VEL 1						
1.	(d)	2.	(b)	3.	(b)	4.	(b)	5.	(b)	6.	(d)	7.	(b)	8.	(a)
9.	(b)	10.	(d)	11.	(c)	12.	(c)	13.	(b)	14.	(c)	15.	(c)	16.	(b)
17.	(d)	18.	(d)	19.	(c)	20.	(a)	21.	(c)	22.	(b)	23.	(a)	24.	(a)
25.	(c)	26.	(c)	27.	(c)	28.	(c)	29.	(a)	30.	(c)	31.	(b)	32.	(b)
33.	(c)	34.	(d)	35.	(b)	36.	(b)	37.	(a)	38.	(c)	39.	(b)	40.	(a)
41.	(d)	42.	(c)	43.	(c)	44.	(c)	45.	(c)	46.	(a)	47.	(a)	48.	(d)
49.	(c)	50.	(d)												



1. Comprehension

For the given question (1, 2, 3), consider the following reaction.

人				
		light		
1 1	$+ X_2 -$	\rightarrow	monohalogenation	product
	4			A 1

- A. Light is involved in which step of the reaction:
 - (a) Initiation only

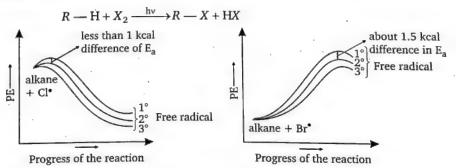
(b) Termination only

(c) Propagation only

- (d) Propagation and Termination
- Which halogen will give the best yield of a single monohalogenation product?
- (b) Cl₂
- (c) Br₂
- (d) I_2
- **C.** How many monohalo derivatives are possible (excluding stereoisomers)?
 - (a) 3
- (b) 4
- (c) 5
- (d) 6

2. Comprehension

Halogenation is a substitution reaction, where halogen replaces one or more hydrogens of hydrocarbon.



Chlorination is exothermic and transition state resembles with products

Bromination is endothermic and transition state resembles with products

Chlorine free radical make 1°, 2°, 3° radicals with almost equal ease, whereas bromine free radicals have a clear preference for the formation of tertiary free radicals. So, bromine is less reactive, and more selective whereas chlorine is less selective and more reactive.

The relative rate of abstraction of hydrogen by Br $^{\bullet}$ is $3^{\circ} > 2^{\circ} > 1^{\circ}$

The relative rate of abstraction of hydrogen by Cl^{\bullet} is : $3^{\circ} > 2^{\circ} > 1^{\circ}$

$$3^{\circ} > 2^{\circ} > 1^{\circ}$$

(5) (3.8) (1)

Consider the above argument and answer A to G:

- 1-halo-2,3-dimethyl butane will be obtained in better yields, if halogen is :
- (a) Br₂ В.

Above product will obtained in better yield if *X* is

- (a) Cl₂
- (b) I₂
- (c) Br₂

(c) I₂

(d) Can't be predicted

(d) Can't be predicted

 CH_3 —CH— CH_3 — Cl_2/hv Product C.

Major product in the above reaction is:

- $\begin{array}{c} \text{CH}_{3} \\ \text{(a) CH}_{3} \text{CH}_{2} \text{Cl} \\ \text{(b) CH}_{3} \text{CH}_{2} \text{CH}_{3} \\ \text{Cl} \\ \text{(c) CH}_{3} \text{CH}_{2} \text{CH}_{2} \text{Cl} \\ \text{(d) CH}_{3} \text{CH}_{2} \text{CH}_{2} \text{CH}_{3} \\ \text{Cl} \\ \end{array}$

- Which of the following will give five monochloro products, when allowed to react with Cl, in D. presence of sun light (excluding stereoisomers)?
 - (a) *n*-pentane
- (b) Iso-pentane (c) 2-methyl-pentane (d) 3-methyl pentane
- 2, Bromo-2, 5, 5 trimethyl pentane (x%)

What is the value of x (% yield of product)?

- (a) 18 %
- (b) 82 %
- (c) 90 %
- What would be the product ratio x/y in the chlorination of propane if all the hydrogen were F. abstracted at equal rate?

- How many dichloro products (including stereoisomers) will be formed when R-2-chloropentane reacts with Cl₂ in presence of UV radiation?
 - (a) 5
- (c) 7
- (d) 8

3.
$$Na \longrightarrow (A)$$
; Product (A) is:

(a)
$$(b)$$
 (c) (d) (d)

5. Ph—CH₂—C—OH
$$\xrightarrow{\text{(1) NaOH, CaO, } \Delta}$$
 (A)

Product (A) is:

(d)

6. Match the column I with column II and with column III.

	Column (I)	- 2,	Column (II)		Column (III)	
		Me	ono-chloro products	Me	onochloro products	
Compound			(excluding stereoisomerism)	(including stereoisomerism)		
(a)		(p)	1	(w)	1	
(b)	$\begin{array}{c} \operatorname{CH}_3 - \operatorname{CH} - \operatorname{CH}_2 - \operatorname{CH}_3 \\ \\ \operatorname{CH}_3 \end{array}$	(q)	2	(x)	3	
(c)	$\begin{array}{c c} \operatorname{CH_3CH_3} \\ \mid & \mid \\ \operatorname{CH_3-C-C-CH_3} \\ \mid & \mid \\ \operatorname{CH_3CH_3} \end{array}$	(r)	3	(y)	5	
(d)	CH ₃ — CH ₂ — CH ₂ — CH ₃	(s)	4	(z)	6	

A. R-2-chloropentane C: Optically active di-chloro products (P)
 B. C: Optically active dichloro products (Q)
 C. R-2-chlorobutane C: Optically active di-chloroproducts (R)

Sum P + Q + R is:

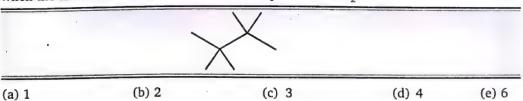
8. Match the column I and II.

	Column (I)		Column (II)	
	Reaction	Type of Reaction		
(a)	CH_3	(p)	Meso compound	
(ъ)	CH_3 CH_3	(q)	Diastereomers	
(c)	CH ₃ O CH ₂	(r)	Racemic	
(d)	H H H H	(s)	Optically inactive due to absence of chiral center	

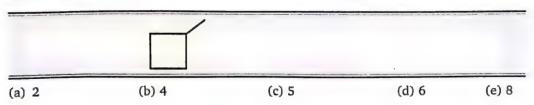
9. Match the column:

	Column (1)		Column (II)	
	Reaction	Product		
(a)	$ \begin{array}{c} \text{CH}_{3} \\ \hline $	(p)	CH ₃	
(b)	$ \begin{array}{c} \text{CH}_{3} \\ \hline $	(q)	CH ₃ DH	
(c)	$ \begin{array}{c} \text{CH}_{3} \\ \hline $	(r)	CH ₃	
(d)	$ \begin{array}{c} \text{CH}_{3} \\ \xrightarrow{\text{(1) BH}_{3}: \text{THF}} \\ \xrightarrow{\text{(2) CH}_{3}\text{CO}_{2}\text{D}} \end{array} $	(s)	CH ₃	

10. How many distinct monochlorinated products, (including stereoisomers) may be obtained when the alkane shown below is heated in the presence of Cl_2 ?



11. How many distinct monochlorinated products, (including stereoisomers) may be obtained when the alkane shown below is heated in the presence of Cl₂?



12. Match the column:

E	Column (1)		Column (II)			
	Wurtz reaction		Number of dimerization product			
(a)	$CH_3 - Cl \xrightarrow{Na} dry ether$	(p)	5			
(b)	$CH_3 - Cl + CH_3 - CH_2 - Cl \xrightarrow{Na}_{dry \text{ ether}}$	(q)	6			
(c)	$\begin{array}{c} \operatorname{CH}_3 - \operatorname{Cl} + \operatorname{CH}_3 - \operatorname{CH}_2 - \operatorname{Cl} \\ + \operatorname{CH}_3 - \operatorname{CH}_2 - \operatorname{CH}_2 - \operatorname{Cl} \xrightarrow{\operatorname{Na}} \\ & \text{dry ether} \end{array}$	(r)	. 3			
(d)	$H_2C = CH - CH = CH - CH_2 - Cl$ + $CH_3 - CH_2 - Cl$ \xrightarrow{Na} $\xrightarrow{dry \text{ ether}}$	(s)	1			

13.
$$H \xrightarrow{CH_3} Cl$$
 $\xrightarrow{Cl_2} CH_2 - CH_2 - CH_3 \xrightarrow{hv} (x)$. $(x) = \text{total number of di-chloro product}$

S-2-chloro hexane

HYDROCARBONS (ALKANES)

189

ANSWERS — LEVEL 2

- 1. A a; B c; C b
- 2. A b; B c; C a; D c; E c; F b; G c
- 3. a, b, c
- **4.** a
- **5.** c
- 6. a-q-x; b-s-z; c-p-w; d-q-x
- 7. P + Q + R = 10
- 8. a-q; b-p; c-r; d-s
- 9. a-p; b-s; c-q; d-r
- **10.** a
- **11.** e
- **12.** a-s; b-r; c-p; d-q
- **13.** 9

4B

HYDROCARBONS (ALKENES)



LEVEL- JE

1. (R)-3-bromocyclopentene (shown below) reacts with Br₂/CCl₄ to form two products, Y and Z, Y is not optically active (does not rotate plane-polarized light). What is the structure of Y?

(a)
$$Br$$
 (b) Br (c) Br (d) Br (e) Br Br Br

(d) All of these

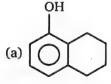
2.
$$A \xrightarrow{\text{2HCl}} Cl$$
 Reactant (A) can be:

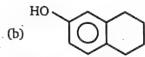
; Major product of the reaction is :

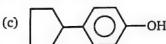
Which of the following products cannot be obtained in ozonolysis of o-xylene?

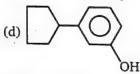
(c)
$$CH_3 - C - C - CH_3$$

→ Major product of the reaction is : 5.









 $CH_2 - CO_2K$ → (A) (Kolbe electrolysis method)

Product (A) of the reaction is:

(a) $CH_3 - CH_3$ (c) $CH_3 - CH = CH_2$

(b) $CH_2 = CH_2$

(d) none of these

 $\xrightarrow{O_3} A \xrightarrow{H_2/Ni} B \xrightarrow{H^+} \Delta$ (C); Product (C) of the reaction is:









8. OH
$$\xrightarrow{Br_2} W$$
. Product W is :

(a) OH

(b) \xrightarrow{Br}

OH

(c) \xrightarrow{Br}

OH

(d) \xrightarrow{Br}

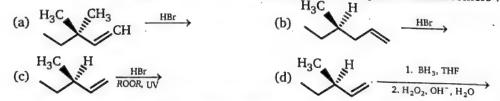
OH

9. The reaction of propene with H_3O^+ will proceed with which of the following intermediates ?

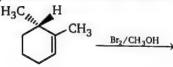
10. Which of the following bromides is the major product of the reaction shown below, assuming that there are no carbocation rearrangement?

$$\begin{array}{c|c} & + & HBr \\ & + & (1 \text{ equivalent}) & \longrightarrow C_{13}H_{17}Br \\ \\ & Br & \\ & (a) & \\ & (b) & \\ & & (c) & Br & \\ & & (d) & \\ & & Br \\ \\ & & \\$$

11. Which of the following reactions results in the formation of a pair of diastereomers?

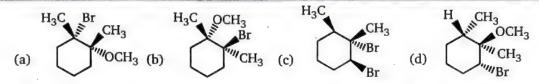


12. What is a likely product of the reaction shown?



HYDROCARBONS (ALKENES)

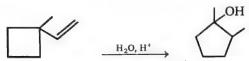
193



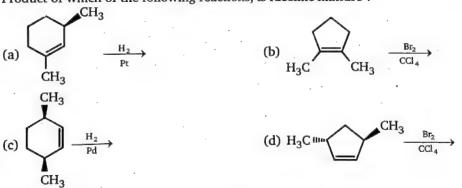
13. Which of the following, when undergoing addition of HBr, will form ONLY a pair of diastereomers?



14. How many transition states and intermediates will be formed during the course of following reaction?



- (a) 3 transition states and 3 intermediates (b) 4 transition states and 3 intermediates
- (c) 3 transition states and 2 intermediates (d) 5 transition states and 4 intermediates
- 15. Product of which of the following reactions, is racemic mixture?

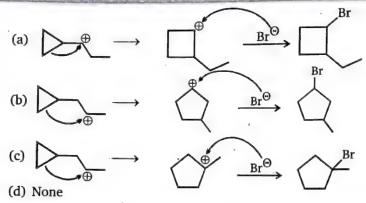


16. The product(s) of the following reaction can best be described as:

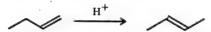
(a) a racemic mixture

- (b) a single enantiomer
- (c) a pair of diasteriomers
- (d) an achiral molecule
- 17. Taking into account the stability of various carbocations and, as well as the rules governing mechanisms of carbocation rearrangements, which reaction is most likely to occur during the given reaction?

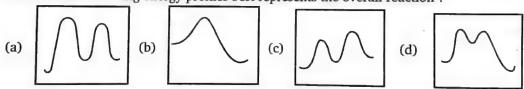
$$H \xrightarrow{Br} \rightarrow ?$$



18. Consider the following reaction in which the intermediate carbocation loses H+ to give the final product?



Which of the following energy profiles best represents the overall reaction?



19. Methyl vinyl ether, $H_2C = CH - OCH_3$, reacts with Br_2/CH_3OH . If methanol is reacting as water would, and if this reaction follows a typical mechanism of electrophilic addition, what would be the expected product?

(a)
$$H_3CO$$
 OCH (b) H_3CO OCH₃ (c) H_3CO OCH₃ (d) H_3CO OCH₃

2, 4-hexadiyne (C₆H₆) is allowed to react with Li in NH₃(liq). The product obtained is 20. treated with 1 equivalent of Cl₂ in CCl₄. Which of the following constitutional isomers are possible products?

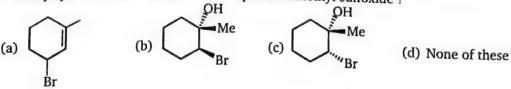
(a) I and II

(b) II and III

(c) I and V

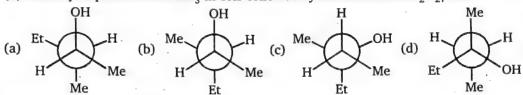
(d) I and III

Which of the following is the best stereochemical representation when reaction between 21. 1-methylcyclohexene and NBS react in aqueous dimethyl sulfoxide?

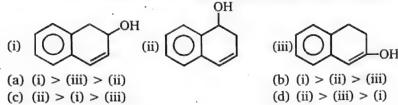


HYDROCARBONS (ALKENES)

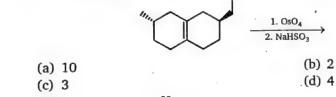
22. Which of the following is among the major products of the reaction of (E)-3-methyl-2-pentene with BH $_3$ in THF followed by the addition of H $_2$ O $_2$ /HO $^-$?



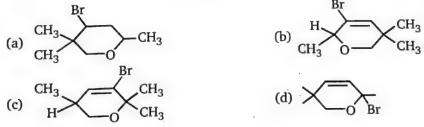
23. Compare rate of dehydration of (i), (ii) and (iii) by conc. H₂SO₄.



24. How many products will be formed in this reaction?

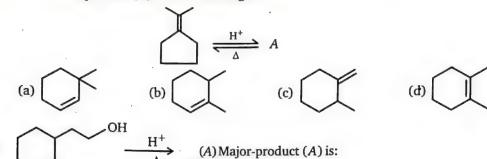


25. C = C = C $C(CH_3)_2CH_2OH \xrightarrow{Br_2} (A)$. Product (A) of the reaction is:



26.
$$CH_3$$
 CH_3
 CH_3

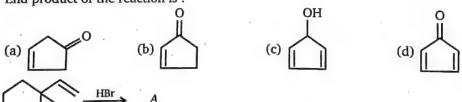
27. Predict the product (*A*) of the following reaction



28.
$$\xrightarrow{H^+} \underset{\text{(Major)}}{\overset{(A) \text{ Major-product } (A) is:}}$$

29. Di-imide
$$(N_2H_4)$$
 is used to reduce double bond of:
(a) $-C = O$ (b) $-C = N$ (c) $-NO_2$ (d) $-CH = CH - CH$
30. $NAHCO_3 \rightarrow (B) \rightarrow (C_5H_8O) \rightarrow (C)$

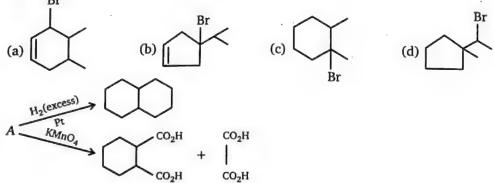
End product of the reaction is:



31. A (Major)

32.

Product (A) is:



33.
$$OsO_4 \longrightarrow X + Y$$

$$12 : 1$$

Product (X) will be:

Product (C) is

35.

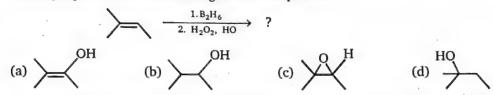
(a)
$${Ph \atop H} \subset C \subset {Ph \atop Ph}$$
 (b) ${Ph \atop H} \subset C \subset {Ph \atop H}$ (c) ${Ph \atop Ph} \subset C \subset CH_2$ (d) $Ph - C \equiv C - Ph$

Product (X) is:

36. Product (P) is:

(d)

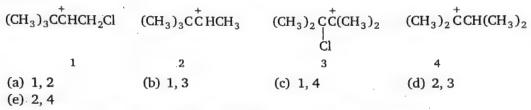
40. The major product of the following reaction sequence is:



41. Which one of the following compounds gives acetone $(CH_3)_2C = O$ as one of the product of its ozonolysis?



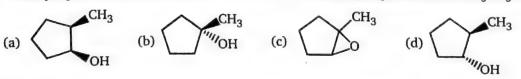
42. Addition of HCl to 3, 3-dimethyl-1-butene yields two products, one of which has a rearranged carbon skeleton. Among the following carbocations, select the possible intermediates in that reaction?



- 43. Conversion of cyclohexene to cyclohexanol can be conveniently achieved by :
 - (a) NaOH + H_2O (b) $Br_2 H_2O$
 - (c) hydroboration, oxidation (d) hydroboration hydrolysis
- 44. Trans-cyclohexane-1,2-diol can be obtained by the reaction of cyclohexene with:
 - (a) $KMnO_4$ (b) OsO_4 (c) peroxy formic acid $/H_3O^+$ (d) SeO_2
- 45. Bromination of (E)-2-butenedioic acid gives
 - (a) (2R, 3S)-2, 3-dibromosuccinic acid
 - (b) (2R, 3R)-2, 3-dibromosuccinic acid(c) a mixture of (2R, 3R) and (2S, 3S)-2, 3-dibromosuccinic acid
 - (d) (2S, 3S)-2, 3-dibromosuccinic acid

(c) $H - C \equiv C - H$

46. The major product formed during the reaction of 1-methyl cyclopentene with CH₃CO₃H is



(d) $CH_2 = CH - CH = CH_2$

49.
$$\underbrace{\begin{array}{c} (1) (CF_3CO_2)_2 Hg, CH_3CH_2OH \\ (2) NaBH_4, HO \end{array}}_{(2) NaBH_4, HO} (P); Product (P) is:$$

$$\underbrace{\begin{array}{c} (2) NaBH_4, HO \\ (2) NaBH_4, HO \end{array}}_{(2) NaBH_4, HO} OCH_2CH_3 (C) OH$$

$$\underbrace{\begin{array}{c} (2) NaBH_4, HO \\ (2) NaBH_4, HO \\ (3) NaBH_4, HO \\ (4) NaBH_4, HO \\ (5) NaBH_4, HO \\ (6) NaBH_4, HO \\ (7) NaBH_4, HO \\ (8) NaBH_4, HO \\ (9) NaBH_4, HO \\ (9) NaBH_4, HO \\ (9) NaBH_4, HO \\ (100\%)$$

50. What is the major product expected from the following reaction?

$$(a) \begin{array}{c} & \underset{H}{\overset{KMnO_4}{\longrightarrow}} \\ & \underset{H}{\overset{OH}{\longrightarrow}} \\ & \underset{OH}{\overset{OH}{\longrightarrow}} \\ & \underset{OH}{\overset{OH}$$

51. $CH_3 - CH = CH_2 \xrightarrow{Br_2/hv} (A)$; Product (A) of the reaction is:

(a)
$$CH_3 - CH - CH_2 - Br$$
 (b) $H_2C = CH - CH_2 - Br$

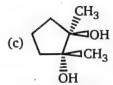
(c) $CH_3 - C = CH_2$ (d) $Br - CH_2 - CH_2 - Br$

$$CH_3 - CH_2 - CH_2$$

53.
$$CH_3$$

$$\xrightarrow{1.0sO_4}$$
 Product of the reaction is:

(a)
$$OH$$
 CH_3 CH_3



54. Which compound is a possible product from addition of Br₂ to 1-butene?

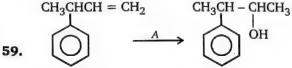
(b) Br

- **55.** Addition of Br_2 to cis-2-butene would give a product which is:
 - (a) achiral

(b) racemic

(c) meso

- (d) optically active
- **56.** Addition of Br₂ to trans-2-butene would give a product which is:
 - (a) achiral
- (b) racemic
- (c) meso
- (d) optically active
- **57.** Addition of OsO₄ to cyclopentene would give a product which is:
 - (a) achiral
- (b) racemic
- (c) meso
- (d) optically active
- **58.** Addition of BH₃ followed by H₂O₂ to trans-2-butene would give a product which is:
 - (a) achiral
- (b) racemic
- (c) meso
- (d) optically active



; Reagent A may be :

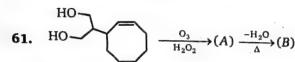
(a) H_2O/H^+

- (b) BH_3 . $THF/H_2O_2 OH^-$
- (c) Hg(OCOCH₃)₂. H₂O/NaBH₄. NaOH
- (d) All are possible
- **60.** The major product of the following reaction is:

$$CH_3 - CH = CH_2 + HBr \xrightarrow{(C_6H_5CO)_2O_2}$$

- (a) CH₃ CH₂ CH₂ Br
- (b) $CH_3CH(Br) CH_3$

- (c) $BrCH_2 CH = CH_2$
- (d) Br



Identify (B):

(a)
$$_{\text{HO}}$$
 (CH₂)₅ - CO₂H (b) $_{\text{HO}}$ (CH₂)₅ - CO₂H (c) $_{\text{HO}}$ (CH₂)₄ - CO₂H

62. Which of the following is a major product of the reaction shown below?

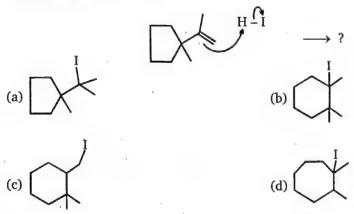
$$(a) \begin{array}{c} CH_3 \\ \hline \\ Br_2 \\ \hline \\ H \end{array} \\ (b) \begin{array}{c} CH_3 \\ \hline \\ UIIIOH \\ \hline \\ H \end{array} \\ (c) \begin{array}{c} CH_3 \\ \hline \\ UIIIBr \\ \hline \\ UIIBR \\ \\ UIIBR \\ \hline \\ UIIB$$

- 63. In methyl alcohol solution, bromine reacts with ethylene (ethene) to yield BrCH₂CH₂OCH₃ in addition to 1, 2-dibromoethane because
 - (a) the methyl alcohol solvates the bromine
 - (b) the ion formed initially may react with ${\rm Br}^-$ or ${\rm CH_3OH}$
 - (c) this is a free radical reaction
 - (d) the reaction follows Markovnikov's rule
- **64.** Which of the following compound was the starting material for the oxidation shown below?

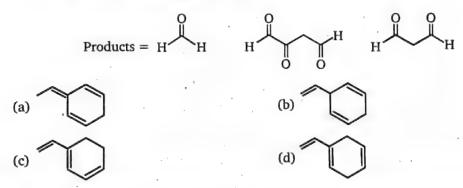
?
$$\xrightarrow{\text{KMnO}_4/\text{H}^+}$$
 $\xrightarrow{\text{HO}}$ $\xrightarrow{\text{OH}}$ $\xrightarrow{\text{OH}}$ $\xrightarrow{\text{OH}}$ $\xrightarrow{\text{OH}}$ $\xrightarrow{\text{CO}_2}$ $\xrightarrow{\text{(a)}}$ $\xrightarrow{\text{(b)}}$ $\xrightarrow{\text{(d)}}$ $\xrightarrow{\text{(d)}}$

65. Which series of reactions will achieve the following transformation?

66. Taking into account the stability of various cycloalkanes and carbocations, as well as the rules governing mechanisms of carbocation rearrangements, what is the most likely product of this reaction?



67. A triene is treated with ozone followed by zinc in acetic acid to give the following three products. What is the structure of the triene?



68. Which of the following compound would yield trialkylborane shown below when treated with $\rm BH_3/THF$?

(a) 2-methylbut-1-ene

(b) 2-methylbut-2-ene

(c) 3-methylbut-1-ene

- (d) 3-methylbut-1-yne
- **69.** If the following compound is treated with Pd/C in excess of hydrogen gas, how many stereoisomers of the product will be obtained?

(a) 1 (c) 3

(b) 2

(d) 4

70. Which is the most precise designation of stereochemistry for the products formed in the electrophilic addition of DBr to 1-methylcyclohexene? ($D = {}^{2}H$, an isotope of hydrogen)

(d) both (a) and (b)

71. Consider the addition of HBr to 3,3-Dimethyl-1-butene shown below. What is the best mechanistic explanation for the formation of the observed product?

- (a) Protonation of the alkene followed by a hydride shift and addition of bromide to the carbocation
- (b) Double bond shift in the alkene following by the protonation and addition of bromide to the carbocation
- (c) Addition of bromide to the alkene followed by a double bond shift and protonation
- (d) Protonation of the alkene followed by a methyl shift and addition of bromide to the carbocation
- **72.** Propene $CH_3CH = CH_2$ can be converted into 1-propanol by oxidation. Indicate which sets of reagents amongst the following is ideal to effect the above conversion?
 - (a) KMnO₄ (alkaline)

(b) Osmium tetroxide (OsO₄/CH₂Cl₂)

(c) B2H6 and alk. H2O2

(d) O_3/Zn

73. Which is the most suitable reagent among the following distinguish compound (3) from the others?

(1) $CH_3C \equiv C - CH_3$

(2) CH₃CH₂ — CH₂ — CH₃

(3) $CH_3CH_2C \equiv CH$

- (4) $CH_3CH = CH_2$
- (a) Bromine in carbon tetrachloride
- (b) Bromine in acetic acid solution

(c) Alk. KMnO₄

(d) Ammonical silver nitrate

74. The principal organic product formed in the reaction given below is:

$$CH_2 = CH(CH_2)_8COOH + HBr \xrightarrow{peroxide}$$

(a) CH₃ — CHBr(CH₂)₈COOH

(b) $CH_2 = CH(CH_2)_8 COBr$

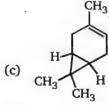
(c) CH₂BrCH₂(CH₂)₈COOH

(d) $CH_2 = CH(CH_2)_7 CHBrCOOH$

HYDROCARBONS (ALKENES)

205

- 75. When 2-butyne is treated with Pd BaSO₄; the product formed will be:
 - (a) cis-2-butene
- (b) trans-2-butene
- (c) 1-butene
- (d) 2-hydroxy butane
- 76. In the reaction, $CH_3C = C CH_3 \xrightarrow{(i) X} CH_3 C C CH_3$, X is:
 - (a) HNO₃
- (b) O₂
- (c) O:
- (d) KMnO
- 77. Which of the following alkene on catalytic hydrogenation given cis and trans-isomer?
 - (a) $H_2C = \bigcirc CH_3$
- (b) H₃C—CH₃



- (d) all of these
- **78.** In the reaction of hydrogen bromide with an alkene (in the absence of peroxides), the first step of the reaction is the to the alkene.
 - (a) fast addition of an electrophilic
- (b) slow addition of an electrophile
- (c) fast addition of a nucleophilic
- (d) slow addition of a nucleophile
- 79. Which of the following alcohols cannot be prepared from hydration of an alkene?
 - (a) 0

(p) OH

- **80.** Which of the species shown below is the most stable form of the intermediate in the electrophilic addition of Cl₂ in water to cyclohexene to form a halohydrin?
 - (a) HHOTH

(c) +

- (d) H
- **81.** The reaction, $(CH_3)_2C = CH_2 + Br^* \longrightarrow (CH_3)_2C CH_2Br$

is an example of a/an step in a radical chain reaction.

(a) initiation

(b) termination

(c) propagation

(d) heterolytic cleavage

- **82.** Which of the following most accurately describes the first step in the reaction of hydrogen chloride with 1-butene?
 - (a) Cl-H +Cl•
 - (b) Cl-H + Cl-
 - (c) Cl-H +Cl-
 - (d) $H CI \longrightarrow CI \longrightarrow +H$
- **83.** Which of the following best describes the flow of electrons in the acid-catalyzed dimerization of $(CH_3)_2C = CH_2$?

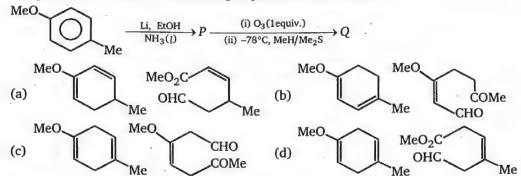
(a)
$$H_{3}C$$
 CH_{3} $H_{2}C = C$ CH_{3} (b) $H_{3}C$ CH_{2} $H_{2}C = CH_{2}$ CH_{3} $CC = CH_{2}$ $CC = CH_{2}$

84. Hydroboration of 1-methylcyclopentene using B₂D₆, followed by treatment with alkaline hydrogen peroxide, gives

The correct statements with respect to the above pair of reactions are that

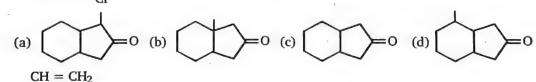
- (I) the reactions are stereospecific
- (II) (X) is erythro and (Y) is threoisomer
- (III) (X) is threo and (Y) is erythro isomer
- (IV) each of (P) and (Q) gives a mixture of (X) and (Y)
- (a) I and II
- (b) I and III
- (c) I and IV
- (d) II and IV

86. The products P and Q in the following sequence of reactions, are



- 87. 4-Pentenoic acid when treated with I_2 and NaHCO₃ gives :
 - (a) 4, 5-diiodopentanoic acid
- (b) 5-iodomethyl-dihydrofuran-2-one
- (c) 5-iodo-tetrahydropyran-2-one
- (d) 4-pentenolyiodide

88.
$$\xrightarrow{\text{H}_2\text{SO}_4, \ 0^\circ\text{C}} \xrightarrow{\text{H}_2\text{O}} (A) \xrightarrow{-\text{HCl}} (B)$$
; Product (B) of the reaction is:



$$\xrightarrow{\text{Br}_2} (A) \xrightarrow{\text{(i) alc.KOH}} (B) \xrightarrow{\text{(i) NaNH}_2} (C), \text{ Product (C) is :}$$

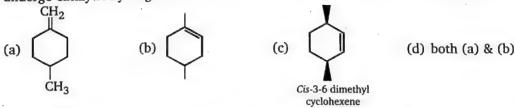
(Styrene)

(a) $Ph - C \equiv CNa$

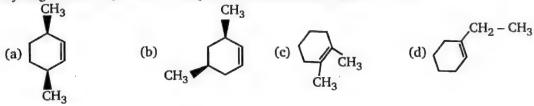
(b) $Ph - CH_2 - C \equiv CH$

(c) $Ph - C \equiv C - CH_3$

- (d) $Ph CH = C = CH_2$
- **90.** Which of the following will give a mixture of *cis* and *trans*-1,4-dimethyl cyclohexane, when undergo catalytic hydrogenation?



91. An optically active compound A with molecular formula C_8H_{14} undergoes catalytic hydrogenation to give meso compound, the structure of (A) is:



92.
$$CH_3 - CH_2 C = CCH_2 - CH_3 + HBr \xrightarrow{R_2O_2 \text{(Per-oxide)}} Products$$

How many products will be formed in above reaction?

(a) 2

(b) 4

(c) 3

(d) 6

93.
$$CH_3$$
 $C = C$ CH_3 Product of the reacion is:

(a) Racemic

(b) Diastereomers

(c) Meso

(d) Pure enantiomers

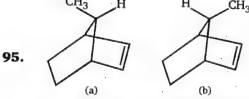
94. cis-2-butene $\xrightarrow{\text{HBr}}$ product; Product of the reaction is:

(a) Racemic

(b) Diastereomer

(c) Meso

(d) E and Z isomer



Rate of reaction towards reduction using (H2/Pt):

(a) a > b

(b) a = b

(c) b > a

(d) Reduction of given molecule is not possible

96.
$$R' \rightarrow C \rightarrow R \xrightarrow{CH_3-S-CH_3} Product A + CH_3 - S - CH_3$$

Product A of the above reaction is:

(b) R' - CHO

(c) $R - CO_2H$

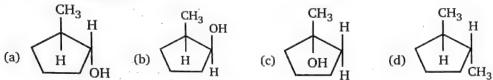
(d) both (a) and (b)

97.
$$CH_3 \longrightarrow C = C \xrightarrow{CH_3} CH_3 \xrightarrow{Cl} CH_2Cl_2 \longrightarrow Product; Product is :$$

MCPBA --- Metachloroperbenzoic acid

(a)
$$\begin{array}{c} CH_3 \\ H \end{array}$$
 $\begin{array}{c} C-C \\ CH_3 \end{array}$ (b) $\begin{array}{c} CH_3 \\ H \end{array}$ $\begin{array}{c} C-C \\ CH_3 \end{array}$ (c) $\begin{array}{c} CH_3 \\ C-C \end{array}$ $\begin{array}{c} CH_3 \\ C-C \end{array}$ $\begin{array}{c} CH_3 \\ C-C \end{array}$ $\begin{array}{c} CH_3 \\ C-C \end{array}$

98.
$$(1) \xrightarrow{\text{BH}_3; \text{THF}} (A) ; \text{ Product of the reaction is :}$$



- **99.** $CH_3 CH = CH_2 \xrightarrow{(1) \text{ THF : BD}_3} (A)$; Product (A) of the above reaction is:
 - (a) CH_3 CHD CH_2D (b) CH_3 — CHT — CH_2T (c) CH_3 — CHD — CH_2T (d) CH_3 — CHT — CH_2D
- **100.** Optically active isomer (A) of (C_5H_9Cl) on treatment with one mole of H_2 gives an optically inactive compound (B) compound (A) will be:
 - (a) $CH_3 CH CH = CH_2$ (b) $CI CH CH = CH CH_3$ (c) $CH_3 CH CH_2 CH = CH_2$ (d) $CH_3 CH_2 CH CH = CH_2$
- 101. An organic compound C_4H_6 on ozonolysis give HCHO, CO_2 , CH_3CHO . Compound will be:

 (a) $H_2C = CH CH = CH_2$ (b) $CH_3 CH = C = CH_2$

(a)
$$CH_3OH$$
 (b) CH_3OH (c) CH_2-OH (d) CH_2-OH (d) CH_3-OH (d) CH_3-OH (d) CH_3-OH (e) CH_3-OH (f) CH_3-OH (f)

Product (C) in the above reactions is:

(a)
$$CH_3 - C - Br$$

$$CH_3$$

$$\begin{array}{c} \operatorname{CH}_3 \\ | \\ \operatorname{C} - \operatorname{Br} \\ | \\ \operatorname{CH}_3 \\ \operatorname{CH}_3 \\ \operatorname{CH}_3 \\ \end{array}$$

(c)
$$CH_3 - CH - H$$

 $CH_2 - Br$

104.
$$CH_3 - C = CH_2 + (CH_3)_2 CHCH_3 \xrightarrow{HF} C_8 H_{18}(A)$$

Unknown (A) in the above reaction is:

- (a) 2, 2, 3-trimethyl pentane
- (b) 2, 2, 4-trimethyl pentane
- (c) 2, 2-dimethyl hexane
- (d) n-octane

105.
$$\xrightarrow{Br_2} HBr + (P) \xrightarrow{MeOH} (Q)$$
; Product (Q) is:

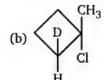
106.
$$\underbrace{\begin{array}{c} H^{+} \\ OH \end{array}}_{OH} \xrightarrow{\Delta} (A) \xrightarrow{\text{cold dil.}}_{KMnO_{4}} (B) \xrightarrow{CrO_{3}} (C)$$

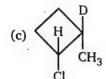
Product (C) of the reaction is:

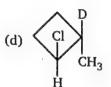
107. What is the major product expected from the following reaction?

$$CH_3 \xrightarrow{D-Cl} Product$$

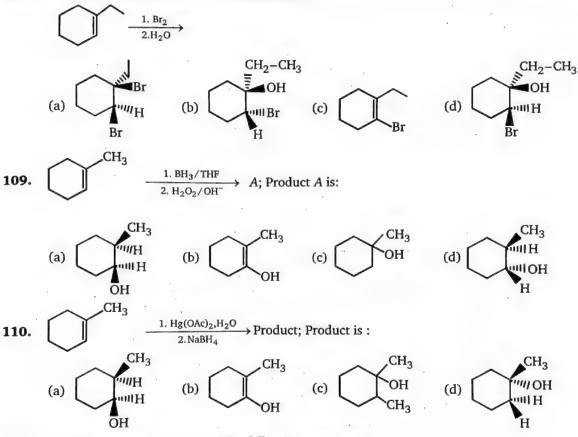




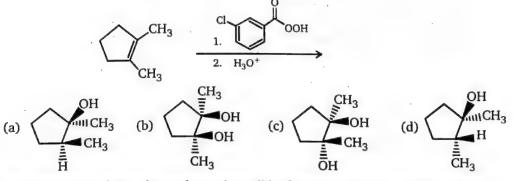




108. Choose the correct product of this reaction:



111. Choose the correct product of the following reactions:



112. How many stereoisomeric tetrabromides will be formed in the following reaction?

113. How many stereoisomeric pentabromides will be formed in the following reaction?

$$\begin{array}{c|c}
Br & 2Br_2 \\
\hline
CCl_4
\end{array}$$

- (a) 2
- (c) 4

- (b) 3
- (d) None of these

114. $\xrightarrow{\text{HCl}} (A) \xrightarrow{\text{EtONa}} (Z)$ (major)

Identify (Z) in the above sequence of reactions:

(a) /

(b) /

(c) O

(d) CH₃

115.
$$CH_3 - CH - CO_2K \xrightarrow{electrolysis} (A)$$
 (Major)
 $CH_3 - CH - CO_2K$

Major product (A) of the above reaction:

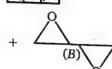
(a) /

(b)

(c) /

(d) i

116.
$$CH = CH_2 \xrightarrow{CF_3CO_3H}$$



(only one enantiomer is taken)

Which of the following statement is correct about A and B?

- (a) A and B are mixture of diastereomers
- (b) A and B are mixture of enantiomers
- (c) A and B are optically active
- (d) B is racemic mixture

117.
$$CH_3O$$

$$\longrightarrow H$$

$$\longrightarrow HOO$$
NaBH₄ $\longrightarrow A$

$$\longrightarrow A$$

$$\longrightarrow B$$

$$\longrightarrow H_2O$$
(C) (one of the product)

Identify the product (C):

118.

O O
$$CH_2 - OH$$

(a) $CH_3 - C - C - O - CH_3$

(b) $CH - OH$
 $CH_2 - OH$
 $CH_2 - OH$

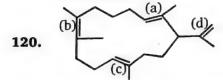
(c) $CH - OH$
 $CH_2 - OH$

(d) $CHOH$
 CHO
 CHO
 CHO
 CHO
 CHO
 CHO
 CHO
 CHO

Product (Y) of the above reaction is:

(a)
$$\bigcirc$$
 (b) \bigcirc CH = CH - CH₂ - OH (c) \bigcirc CH = CH - CH₂ - OH

- In the reaction $Me C \equiv C Et$ \rightarrow (Q); then Q is: 119.
 - (a) A pure compound which is optically inactive due to internal compensation
 - (b) A binary mixture which is optically inactive due to external compensation
 - (c) A binary mixture which is optically active
 - (d) A pure compound which is optically inactive due to absence of chiral centre



Which $(\pi$ -bond) will reduce first, when above compound undergoes catalytic hydrogenation?

(a) a

(b) b

(c) c

(d) d

Compound A, which is a degradation product of the antibiotic vermiculine has following structure 121.

$$CH_{3} \xrightarrow{C} CH \xrightarrow{CH_{2}} CH_{2} \xrightarrow{CH_{2}} CH = CH - CH_{3}$$

$$CH_{3} \xrightarrow{CH_{2}} CH_{2} \xrightarrow{CH_{2}} CH = CH - CH_{3}$$

$$CH_{3} \xrightarrow{CH_{2}} CH_{3} \xrightarrow{CH_{2}} CH_{3} \xrightarrow{CH_{2}} CH_{3}$$

$$CH_{3} \xrightarrow{CH_{2}} CH_{3} \xrightarrow{CH_{2}} CH_{3} \xrightarrow{CH_{2}} CH_{3} \xrightarrow{CH_{2}} CH_{3}$$

$$CH_{3} \xrightarrow{CH_{3}} CH_{3} \xrightarrow{CH_{3}} CH_{3} \xrightarrow{CH_{2}} CH_{3} \xrightarrow{CH_{2}} CH_{3} \xrightarrow{CH_{2}} CH_{3}$$

$$CH_{3} \xrightarrow{CH_{3}} CH_{3} \xrightarrow{CH_{3}} CH_{3} \xrightarrow{CH_{3}} CH_{3} \xrightarrow{CH_{2}} CH_{3} \xrightarrow{CH_{2}} CH_{3} \xrightarrow{CH_{2}} CH_{3} \xrightarrow{CH_{3}} CH_{3} CH_{3} \xrightarrow{CH_{3}} CH_{3} \xrightarrow{CH_{3}} CH_{3} CH_{3}$$

(a)
$$CH_{2}$$
 CH_{2} CH_{3} CH_{2} CH_{2} CH_{2} CH_{3} CH_{2} CH_{2} CH_{3} CH_{3} CH_{3} CH_{2} CH_{2} CH_{3} CH_{3}

(d) None of these

Reagent (A) and (B) in above reaction are:

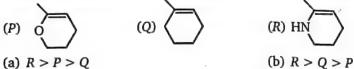
(a)
$$A = RCO_3H$$
, $B = H_2O_2$

(b)
$$A = RCO_3H$$
, $B = HIO_4$

(c)
$$A = RCO_3H, B = O_3$$

(d)
$$A = O_3, B = RCO_3H$$

Rank the following in the increasing order of rate of reaction with HBr. 123.



(b)
$$K > Q > P$$

(c)
$$P > R > S$$

(d)
$$P > S > R$$

Select the reaction(s) that would result in the formation of 2-bromopropane. 124.

(I)
$$CH_3CH = CH_2 + HBr \xrightarrow{peroxide}$$

(II)
$$CH_3CH = CH_2 + HBr \xrightarrow{CCl_4}$$

(III)
$$CH_3CH_2CH_3 + Br_2 \xrightarrow{hv}$$

(IV)
$$CH_3CH = CH_2 + Br_2 \xrightarrow{CCl_4}$$

Which of the following reactions generates the major product? Ignore stereoisomerism. 125.

$$(a) \bigcirc + HBr \longrightarrow \bigcirc_{B_1}$$

HYDROCARBONS (ALKENES)

215

(b)
$$\underbrace{ \begin{array}{c} \text{(1) Hg(OAc)}_2, \text{H}_2\text{O}, \text{THF}} \\ \text{(2) NaBH}_4 \end{array} }_{\text{(2) NaBH}_4} \text{OH}$$
(c)
$$\underbrace{ \begin{array}{c} \text{(1) BH}_3 \\ \text{(2) OH}^-, \text{H}_2\text{O}_2, \text{H}_2\text{O} \end{array} }_{\text{OH}} \text{OH}$$
(d)
$$\underbrace{ \begin{array}{c} \text{H}_2\text{O}, \text{H}_2\text{SO}_4 \\ \text{CCI}_4 \end{array} }_{\text{Br}} \text{Br}$$

126. In the given selective hydrogenation which combination is incorrect?

(a)
$$\xrightarrow{\text{H}_2}$$
 $\xrightarrow{\text{W.C.}}$ (W.C. = Wilkinsons catalyst)

(b)
$$\xrightarrow{\text{H}_2}$$
 (W.C. = Wilkinsons catalyst)

(c)
$$\frac{H_2}{W,C}$$

(d)
$$\xrightarrow{\text{H}_2}$$
 CH_2 \rightarrow $\text{CH} = \text{CH} - \text{CH}_3$

127.
$$(A) \xrightarrow{\text{Na/NH}_3(l)} (B) \xrightarrow{\text{MCPBA}} (C)$$

Compound (C) in above sequence of reaction is:

(a)
$$\bigcirc$$
 $\stackrel{H}{\bigcirc}$ $\stackrel{H}{\bigcirc}$ (b) \bigcirc $\stackrel{H}{\bigcirc}$ $\stackrel{H}{\bigcirc}$

128.
$$\xrightarrow{\text{HBr}} (A);$$

$$\text{CH}_3 \xrightarrow{R_2 \text{O}_2/h\nu} (A);$$

Major product (A) is:

129. In the reaction given below, the product would be:

$$CH_3 - CH = CH - CH_3 \xrightarrow{H_3O^+} CH_3 - CH_2 - CH - CH_3$$

- (a) a mixture of diastereomers
- (c) optically pure enantiomer
- (b) optically active
- (d) a racemic mixture

130. Surprisingly, the reaction shown below goes through classical carbocation. What is the major product of this reaction?

- (a) trans-1, 3-dibromocyclohexane
- (b) cis-1, 3-dibromocyclohexane
- (c) trans-1, 2-dibromocyclohexane
- (d) cis-1, 2-dibromocyclohexane
- **131.** The major product of the reaction given below is:

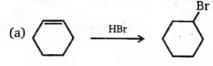
$$OH \xrightarrow{Br_2} ?$$

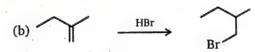
(vi) O CH₂Br

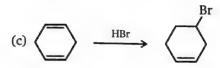
- (a) (i) and (ii)
- (c) (v) and (vi)

- (b) (iii) and (iv)
- (d) none of these

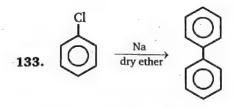
132. Which reaction will occur at the fastest rate?











Above reaction is known as:

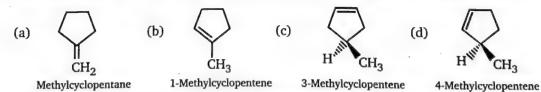
- (a) Wurtz reaction
- (c) Fittig reaction

- (b) Wurtz fittig reaction
- (d) Kolbe electrolysis

134.
$$CH_3$$
— CH_2 — C — H — $Red P + HI$ A

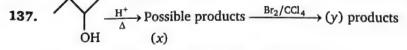
Product A is:

- (a) propane
- (b) propanol
- (c) prapanoic acid
- (d) propene
- 135. Which of the following compound give diastereomers when treated with Br₂ in CCl₄?



- **136.** A mixture of C₂H₆, C₂H₄ and C₂H₂ is bubbled through alkaline solution of copper (1) chloride, contained in Woulf's bottle. The gas coming out is:
 - (a) original mixture

- (b) C_2H_6
- (c) C₂H₆ and C₂H₄ mixture
- (d) C2H4 and C2H2



The number of possible products for x and y is :

(a) 2, 4

(b) 3, 5

(c) 3, 6

(d) 3, 4

- **138.** Select the incorrect statement :
 - (a) Bromine is more selective and less reactive
 - (b) Chlorine is less selective and more reactive
 - (c) Benzyl free radical is more stable than 2° free radical
 - (d) Vinyl free radical more stable than allyl free radical
- **139.** Which of the following compound does not evolve CO₂ gas, when undergo oxidative ozonolysis?



(c)
$$H_2C = CH - CH = CH_2$$

140. *cis*-3-hexene $\xrightarrow{(a)}$ meso 3,4-hexanediol

trans-3-hexene $\xrightarrow{(b)}$ meso 3,4-hexanediol.

Choose pair of reagent (a, b) for above conversions.

(a) Cold KMnO₄, OsO₄

- (b) Cold KMnO₄, RCO₃H/H₃O[⊕]
- (c) RCO₃H/H₃O[⊕], cold KMnO₄
- (d) None of these

141.
$$\underbrace{ \begin{array}{c} \text{Na} \\ \text{Liq. NH}_3 \end{array}}_{\text{Liq. NH}_3} (A) \underbrace{ \begin{array}{c} \text{O}_3 \\ \text{Zn} \end{array}}_{\text{C}} (B) \underbrace{ \begin{array}{c} \text{Ph}_3 \text{P} = \text{CH}_2(2\text{mole}) \\ \text{C} \end{array}}_{\text{Ph}_3} (C)$$

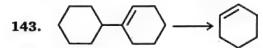
Product (C) of the above reaction is:

(a) 1,3-hexadiene

(b) 1,4-pentadiene

(c) 1,3-butadiene

- (d) 1,3-heptadiene
- **142.** How many carbon-hydrogen bond orbitals are available for overlap with the vacant *p*-orbital in ethyl carbocation ?
 - (a) 0
- (b) 3
- (c) 5
- (d) 6



To achieve above conversion, the reagents used will be:

(a) O_3/H_2O_2 , HO^-/Δ

- (b) HBr, alc. KOH, O_3 , LiAlH₄, H^+/Δ
- (c) HBr, t-BuOK, O_3 , KMn O_4 , Δ
- (d) HCl, KMnO₄ (cold), H^+/Δ

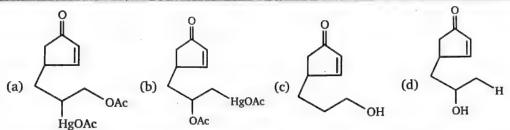




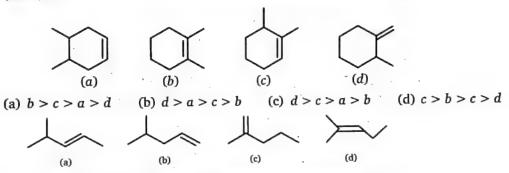
 $\xrightarrow{\text{Hg(OAc)}_2} X$ (major); Product (X) is:

HYDROCARBONS (ALKENES)

219



145. Decreasing order of heat evolved upon catalytic hydrogenation of given reactants with a H₂ (Pd/C) is :



The correct order of heat of hydrogenation of given molecules is :

(a)
$$d > c > a > b$$

146.

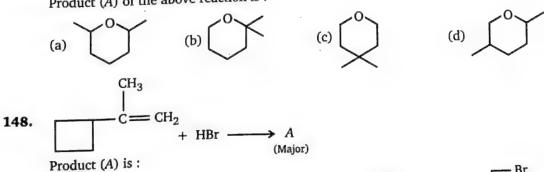
(b)
$$d > c > b > a$$

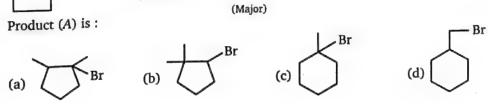
(c)
$$b > a > c > d$$

(d)
$$d > a > c > b$$

147. OH
$$\frac{1. \text{ Hg(OAc)}_2}{2. \text{ NaBH}_4} A$$

Product (A) of the above reaction is:





149.
$$\xrightarrow{\text{HBr}}$$
 Product; Comment upon optical activity of the product.

- (a) Racemic mixture
- (b) Diastereomers
- (c) Meso
- (d) Optically inactive due to absence of chiral center

$$CH_3 - C = CH - CH_3 \xrightarrow{Hg(OAc)_2/EtOH} (A)$$

Product (A) of the above reaction is:

(a)
$$CH_3$$
 CH_3 (b) CH_3 CH_3

(b)
$$CH_3$$
— C — CH — CH_3
OEt HgOAc

(c)
$$CH_3$$
 CH_3 CH_3 (d) CH_3 CH_3

151.
$$Me_2CH \longrightarrow CH \longrightarrow Me \xrightarrow{Al_2O_3} (A) \xrightarrow{(i) HI} (B)$$

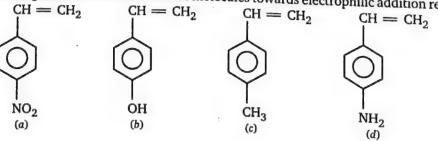
OH

Product (B) of above reaction:

(a) Me₂C(OH)CH₂Me

- (d) $HO CH_2 (CH_2)Me$
- In which of the following reaction, Markownikoff's rule is violated?

 - (a) $CH_3 O CH = CH_2 \xrightarrow{HBr}$ (b) $CH_3 NH CH = CH_2 \xrightarrow{HBr}$ (c) $CH_3 S CH = CH_2 \xrightarrow{HBr}$ (d) $O_2N CH = CH_2 \xrightarrow{HBr}$
- Decreasing order of rate of reaction of molecules towards electrophilic addition reaction is: 153.



(a) a > b > c > d

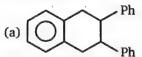
(b) b > c > a > d

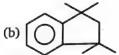
(c) d > b > c > a

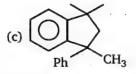
(d) b > d > c > a

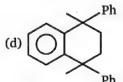
 $2CH_3 - C = CH_2 -$ Ph

Product (A) is:









155. CH₂ -- OH

$$\xrightarrow{H^+} (A) \xrightarrow{\text{ROH}} (B)$$

$$(\text{major})$$

Product (B) of the above reaction is:

(d)
$$\bigcirc$$
 O—OR

Which of the following compounds gives the same carbocation on ionization? 156.



(a) 1 and 3

(b) 2 and 4

(c) 1 and 2

(d) 1 and 4

For the following reactions the major products are shown: 157.

$$H_2C = CH - CH = CH_2 \xrightarrow{HBr} H_2C = CH - CH - CH_3 \xrightarrow{+ 25^{\circ}C} CH_2CH = CHCH_3$$

$$| \qquad \qquad | \qquad \qquad |$$

$$| \qquad |$$

$$| \qquad |$$

$$| \qquad |$$

$$| \qquad |$$

$$| \qquad |$$

$$| \qquad |$$

These provide an example of $\frac{1}{2}$ control at low temperature and $\frac{2}{2}$ control at higher temperature.

- 1

2

- (a) kinetic
- thermodynamic
- (b) thermodynamic
- kinetic

- (c) kinetic
- kinetic
- (d) thermodynamic thermodynamic

158. What is the product of 1, 4-addition in the reaction shown below?

$$(a) \xrightarrow{HCl}$$

$$(b) \xrightarrow{Cl}$$

$$(c) \xrightarrow{Cl}$$

$$(d) \xrightarrow{Cl}$$

Dehydration of the above compound will give:

(a) meso product

(b) racemic mixture

(c) diastereomer

(d) optically pure enantiomer

160.
$$H = CH_{2} - CH_{3}$$

$$CH = CH_{2}$$

$$CH = CH_{2}$$

What is stereochemistry of product?

(a) Racemic mixture

(b) Optically inactive

(c) Diastereomers

(d) Meso product

161.
$$\equiv$$
 OH $\xrightarrow{\text{H}_2}$ A $\xrightarrow{\text{H}^{\oplus}}$ CH₃

End product formed in the above reaction is:

- (a) Optically active (b) Racemic
- (c) Meso
- (d) Diastereomer
- How many moles of BH3 are needed to react completely with 2 mole of 1-pentene in 162. hydroboration-oxidation reaction?
 - (a) 2 mole

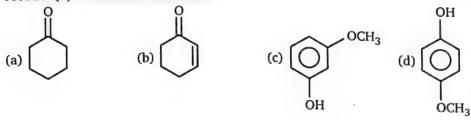
(b) 3 mole

(c) 2/3 mole

(d) 3/2 mole

163.
$$\underbrace{\text{Li}}_{\text{Liq. NH}_3} A \xrightarrow{\text{H}_3\text{O}^+} A$$

Product (B) in the above reaction is:



164.
$$H_2^{14} = CH - CH_3 \xrightarrow{low conc. of Br_2 \ or high temp} (?)$$

Product of the above reaction is:

(a)
$$H_2^{14} = CH - CH_2 - Br$$

(b)
$$H_2C = CH - CH_2 - Br$$

(c)
$$\stackrel{14}{\overset{}{\text{CH}}}_2$$
 — $\stackrel{\text{CH}}{\overset{}{\text{-CH}}}$ — $\stackrel{\text{CH}}{\overset{}{\text{-CH}}}_3$

(d) both (a) and (b)

165. In which of the following reactions 1,3-butadiene will be obtained as a major product?

(a) Br
$$- \text{CH}_2 - \text{CH}_2 - \text{CH}_2 - \text{CH}_2 - \text{Br} \xrightarrow{\text{(CH}_3)_3 \text{COK (2mole)}}$$

(b)
$$HO - CH_2 - CH_2 - CH_2 - CH_2 - OH \xrightarrow{Conc. H_2SO_4}$$

(c) $H_2C = CH - C \equiv CH \xrightarrow{N_{12}B}$

(c)
$$H_2C = CH - C \equiv CH \xrightarrow{H_2(1\text{mole})} N_{1_2B}$$

(d) All of these

166.
$$H_2C = C \xrightarrow{CH_3} \xrightarrow{Cl_2} \xrightarrow{1. H_2O} \xrightarrow{H^+} A$$
; Identify A.

(c)
$$CH_3 - C - CH_2 - CH_3$$

(d)
$$CH_3 - C = CH_2$$

Product (A) is:

(d) d

168.
$$CH_3$$
 d c $NBS \rightarrow A$ (major)

H

 CH_3

Bromination take place at:

(a) a (b) b (c) c

169. Which is incorrect statement about heats of combustion?

(c) Iso-butene > trans-2-butene > 1-butene (d) n-Hexane < n-Heptane < n-Octane

170. Predict the major product of the reaction.

$$\begin{array}{c} \text{CH}_{3} \\ \text{CH}_{2} \\ \text{CH}_{3} \\$$

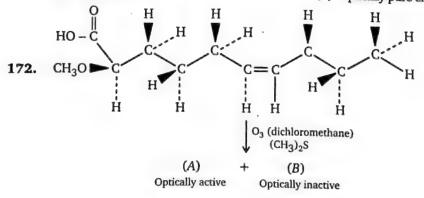
 $cold dil. KMnO_4 \rightarrow Product of the reaction is:$ 171.

- (a) Meso compound

(b) Enantiomeric pair

(c) Diastereomers

(d) Optically pure enantiomer



Product (A) of above reaction is:

(a)
$$CH_3O - CH - CH_2 - CH_2 - CHO$$

 CO_2H CO_2H
(b) $CH_3O - CH_2 - CH - CH_2 - CO_2H$

(c)
$$CH_3O - CH - CH_2 - CH_2 - CO_2H$$

 CO_2H

(d)
$$CH_3O$$
— CH — CH_2

173.
$$H_2$$
 (2-3atm)(1 mole) Products; Comment up on optical activity of products.

- (a) Diastereomers
- (b) Racemic mixture (c) Meso
- (d) Optically pure enantiomer

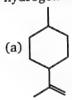
174.
$$+ HBr \longrightarrow Product$$

Addition of a mineral acid to an olefin bond leads to major product, Identify it:

(c)
$$\bigcup_{Br}^{O}$$

175.
$$\frac{H_2(\text{one mole})}{\text{PtO}_2} \text{ Product}$$

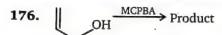
In polyenes that contain differently substituted (C=C) double bonds, it is possible to hydrogenate chemeselectively one (C=C) double bond. Product is:











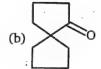
(MCPBA -----> meta-chloro perbenzoic acid)

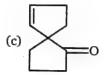
Stereochemistry of the product of above reaction is :

- (a) Meso
- (b) Racemic
- (c) Diastereomers
- (d) Optically inactive due to absence of chiral center.

Identify product (P).



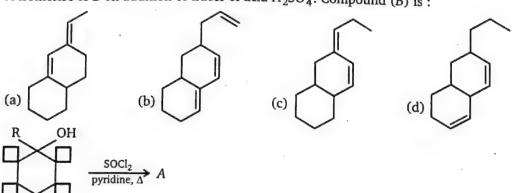




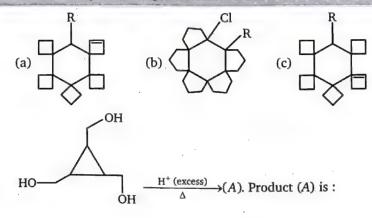


179.

A isomerise to B on addition of traces of acid H_2SO_4 . Compound (B) is:

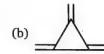


Product (A) of the reaction is:

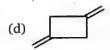




180.

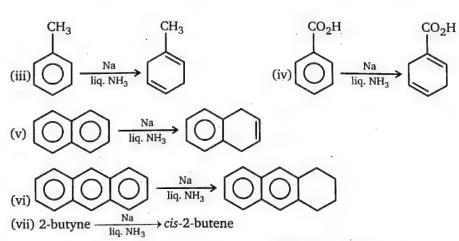






(d) None of these

181. Which of the following reactions do not represent the major product of given Birch reductions?

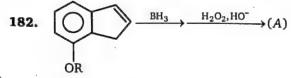


(a) (i), (iii), (vi)

(b) (iv), (vi), (vii)

(c) (iv), (v), (vi)

(d) (i), (ii), (v), (vii)



Product (A) is:

Hint: Think carefully about the relative stabilization of developing positive charge, when the double bond reacts with an electrophile.

183.
$$\underbrace{\qquad \qquad}_{\text{HOCl}} \xrightarrow{\text{HOCl}} (A) \xrightarrow{\text{NaOH,}}_{\text{H2O}} \xrightarrow{\text{70\%}} (B)$$

Correct statement about above reaction is:

(a) A = cis-2-chlorocyclohexanol,

B = cyclohexeneoxide

(b) A = trans-2-chloro cyclohexanol,

B = anti-diol

(c) A = trans-2-chlorocyclohexanol,

B = cyclohexeneoxide

(d) A = cis-2-chlorocyclohexanol,

B = anti-diol

184.
$$\xrightarrow{H^{\oplus}}$$
 Predict the major product:

185.
$$H_2SO_4 \rightarrow (A)$$
; Product (A) is:

0

186.
Br
 $^{82}Br - ^{82}Br \xrightarrow{CCl_4}$ Major product of the reaction is :

187. stereochemistry of the product is: H CH₃

(a) Diastereomers

(b) Racemic mixture

(c) Meso

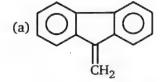
- (d) Pure Enantiomers
- 188. → Product/s obtained is/are:
 - (a) Diastereomers

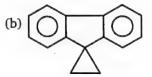
(b) Racemic

(c) Meso

(d) Optically pure enantiomers

189.
$$\xrightarrow{\text{Ph}_3\text{P}=\text{CH}_2} \xrightarrow{\text{Ph}_3\text{P}=\text{CH}_2} (x); \text{ Product } (x) \text{ is :}$$





CH₃

CH₃

CH₃

CH₃

CH₃

CH₃

(A); Product (A) is:

$$\begin{array}{c}
CH_3 \\
CH_3
\end{array}$$
(a) CH₃ - CH₂ - CH₂ - CH₂ - CH₂ - CH₂ - CH₃

(a)
$$CH_3 - CH_2 - CH_2 - CH_2 - CH_2 - CH_3 - CH_3$$

(b)
$$CH_3$$
— CH_2 — CH_2 — CH_2 — $CH = C < CH_3 < CH_3$

(c)
$$CH_3 - CH_2 - CH_2 - CH_2 - CH_2$$

(d)
$$CH_3 - CH_2 - CH - CH_2 - CH_2$$

 CH_3

191.
$$CH_3 - CH = CH - CH_3 \xrightarrow{HBr \\ R_2O_2, \Delta}$$
 (Anti-Markownikoff's addition)

Comment on optical activity of the products:

- (a) Racemic
- (c) Meso

- (b) Diastereomer
- (d) Optically pure enantiomer

192.
$$\xrightarrow{\text{CH}_3-\text{C-O-O-H}} (A); \text{ Product } (A) \text{ is :}$$

193.
$$A$$
(alkene)

Cold dil. KMnO₄
Meso-compound

Alkene (A) will be:

(a) cis-2-pentene

(b) cis-2-hexene

(c) cis-4-octene

(d) trans-2-hexene

194.
$$H \mapsto (A) + PPh_3 \longrightarrow (A) + POPh_3 \mapsto (Major) + ($$

Product (A) is

- (a) trans-2-butane
- (b) cis-2-butene
- (c) 1-butene
- (d) Iso-butene
- **195.** In which of the following reactions, two products will be formed other than phosphonium ylide (POPh₃)

(a)
$$Ph_3P = CH_2 \longrightarrow$$

(b)
$$CH_3CHO + Ph_3P = C - CH_3 \longrightarrow CH_3$$

(c)
$$Ph - C - H + Ph_3P = CH - Ph \longrightarrow$$

(d)
$$H - C - H + Ph_3P = CH - CH_3 \longrightarrow$$

196. To carry out the given conversions, select the correct option:

- (a) $a = \text{Ag}_2\text{O}$,
- $b = \text{Zn/CH}_3\text{CO}_2\text{H}, \quad c = \text{LiAlH}_4$
- (b) $a = H_2O_2$,
- $b = CH_3 S CH_3$, $c = NaBH_4$
- (c) Both (a) and (b)
- (d) None of these
- **197.** The product (A) of given alkoxymercuration de-mercuration is :

$$(a) \xrightarrow{(1)\text{Hg}(O_2\text{CCF}_3)_2, \text{ CH}_3\text{OH}} (A)$$

$$(a) \xrightarrow{(2)\text{NaBH}_4, \text{HO}^-} (C) \xrightarrow{(C)} (C) (C)$$

$$(C) \xrightarrow{(C)} (C) (C)$$

$$(C) \xrightarrow{(C)} (C)$$

198.
$$CH_3 \xrightarrow{C} CH_2 \xrightarrow{HC \equiv CH} \xrightarrow{H^+} \xrightarrow{H_2} \xrightarrow{H_2 \to Al_2O_3} \xrightarrow{\Lambda}$$

End product of the reaction is:

- (a) $H_2C = CH C = CH_2$ CH_3
- (b) $CH_3 CH = CH CH = CH_2$
- (c) $H_2C = CH CH = CH_2$
- (d) $H_2C = CH CH_2 CH = CH_2$
- 199. Major product of the given reaction is:

$$H_2C = CH - CH_2 - I \xrightarrow{HI(excess)} CCI_4 \rightarrow$$

(a) CH₃ — CH— CH₂

- (b) CH₃ CH CH₃
- (c) $CH_3 CH_2 CH_2 I$
- (d) $I CH_2 CdH_2 CH_2 I$

- **200.** The rate constant for a reaction can be increased by \underline{a} the stability of the reactant or by \underline{b} the stability of the transition state. Select the correct choice for a and b.
 - (a) decreasing, decreasing

(b) increasing, decreasing

(c) decreasing, increasing

- (d) increasing, increasing
- **201.** Major product of the given reaction is:

$$H_2C = CH_2 + CH_2$$
 $\xrightarrow{H^+} Product$

$$CH_3$$

|
(d) $H_2C = C - CH_2 - CH_2 - CH_3$

202.
$$+ Ph_3P = CH_2 \longrightarrow (A)$$
Majo

Major product (A) is:

203. In the given reaction, only one alkene undergo preferential oxidation by electrophilic ozone. Identify product (*P*) of the given reaction:

OMe
$$\begin{array}{c}
O_{3} \\
Me_{2}S, -78^{\circ}C \text{ then NaBH}_{4}
\end{array}$$
(P)
$$(>52\%)$$
(a)
$$CO_{2}Me$$
(b)
$$CO_{2}Me$$
OH

(c) $CH_{3} - C - CH_{2} - CH = C - CH_{2} - CHO$
OMe

(d) $CH_3 - CH - CH_2 - CH = C - CH_2 - CH_2 - OH$ OMe

204.
$$CH_2-CH_2-CH = CH_2 \xrightarrow{HCl} (P)$$
; Product (P) is:

 \rightarrow (A); Major product of the reaction is: 205.

206. OMe
$$\xrightarrow{\text{H}_2}$$
 (A) $\xrightarrow{\text{H}_3O^{\circledast}}$ (B)

Product (B) is:

OH
$$\parallel$$
 (a) Ph - CH - CH = CH - CH₂ - C-H

(b)
$$Ph - CH = CH - CHO$$

(c)
$$Ph - (CH = CH)_2 - CHO$$

(d)
$$Ph - (CH = CH)_3 - CHO$$

Isobutene, in the presence of H₂SO₄, forms a mixture of two isomeric alkene (C₈H₁₆). The 207.

(b)
$$CH_3 - CH_3 - CH_2 - CH_2 - CH_2$$

$$CH_3 - CH_2 - CH_2 - CH_2$$

$$CH_3 - CH_3 - CH_3$$

(d)
$$CH_2 = C - CH_2 - CH_2 - CH - CH_3$$

209.

An unknown alkene (A) reacts with 3 mole of H_2 gas in presence of platinum catalyst to form 208. 1-isopropyl-4-methyl cyclohexane. When unknown alkene (A) is ozonized and reduced, following product are obtained

The alkene (A) is:

Product (C) is

210. The following reaction take place in high yields.

$$\begin{array}{c}
CO_2CH_3 \\
& \xrightarrow{Hg(OAc)_2}
\end{array}$$
Product

Use your knowledge of alkene chemistry to predict a product even though you have never seen this reaction before

$$(d) \begin{array}{c} CO_2CH_3 \\ OH \end{array}$$

What is the ratio of glyoxal to pyrualdehyde obtained in the above reaction?

Which of the following product cannot be obtained in above reaction?

(a)
$$H - C - CH_2 - C - H$$

(b)
$$CH_3 - C - CH_2 - C - H$$

213.
$$CH_3 = C \xrightarrow{CH_3} CH_3 + (CH_3)_3 \stackrel{+}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{CH_3} CH_3 \stackrel{-}{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mole})} A + (CH_3)_3 \stackrel{N}{N} = O + H_2O \xrightarrow{OsO_4(10^{-4} \text{ mol$$

2, 3-dimethyl-2-butane (0.025 mole)

(TMAO \rightarrow trimethyl amine -N - oxide) Product (A) is:

(a)
$$CH_3$$
 $C-C$ CH_3 CH_3

$$\begin{array}{c|c} CH_3 CH_3 \\ \mid & \mid \\ (b) CH_3 - C - C - CH_3 \\ \mid & \mid \\ OH OH \\ O \end{array}$$

Product (A) of the reaction is:

(d) None of these

215.
$$OH \xrightarrow{H^+} (A) \atop (major)$$

Product (A) is:

CH₃
$$C = CH_2$$
 CH_3 $C = CH_2$ CH_3 CH_3

Arrange the above in the decreasing order of reactivity towards HBr:

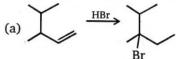
(a) a > b > c

(b) b > a > c

(c) b > c > a

(d) a > c > b

217. Which reaction has the lowest ΔG^{\ddagger} or (Activation-Energy)?



(b)
$$\xrightarrow{\text{HBr}}$$
 $\xrightarrow{\text{Br}}$

(c)
$$\xrightarrow{\text{HBr}}$$
 $\xrightarrow{\text{Br}}$

$$(d) \xrightarrow{HBr} \xrightarrow{Br}$$

218. Which of the following will rearrange?









(a) 1

(b) 1 and 3

(c) All

(d) 1, 2, 4,

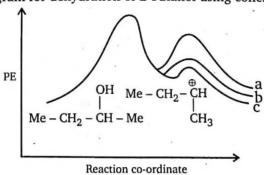
219. Which of the following is most likely to undergo a favorable hydride shift?







220. Energy profile diagram for dehydration of 2-butanol using conc. H_2SO_4 is given below:



Product (b) of above reaction is:

(a) 1-butene

(b) cis-2-butene

(c) trans-2-butene

(d) iso-butene

- 221. How many alkene on catalytic hydrogenation given isopentane as a product?
 - (a) 2

(b) 3

(c) 4

- (d) 5
- 222. Which of the following would not rearrange to a more stable form ?



(b) H

(c) +

- (d) +
- 223. Consider the following reaction.

$$BrCH_2CH_2F + SbF_5 \xrightarrow{SO_2} CH_2 \xrightarrow{CH_2} CH_2 + SbF_6$$

In this reaction SbF5 acts as:

(a) an acid

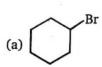
(b) a base

(c) a nucleophile

(d) an electrophile

224.
$$\longrightarrow$$
 $\xrightarrow{\text{Br}_2/\text{hv}}$ Major (X) $\xrightarrow{\text{Alcoholic}}$ $\xrightarrow{\text{KOH}/\Delta}$ Major (Y) $\xrightarrow{\text{H-Br}}$ Major (Z) :

Product (Z) is:



(b) D

(c) Br

(d) B1

225.
$$CH_3-C \equiv C-H \xrightarrow{NaNH_2} CH_3-I \xrightarrow{Li/liq} (B)$$

Relation between (B) and (C) is:

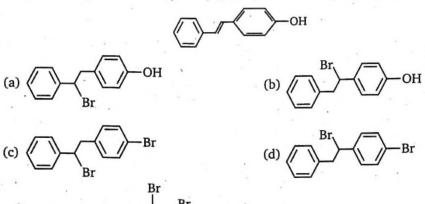
(a) Enantiomer

(b) Diastereomer

(c) Geometrical isomer

(d) Meso

226. The reaction of HBr with the following compound would produce:



- **227.** \longrightarrow + Br₂ \longrightarrow is an example of:
 - (a) Nucleophilic addition
 - (c) Electrophilic addition
 - (e) Free radical substitution
- 228. Olefins can be hydrogenated by:
 - (a) Zinc and HCl
 - (c) Raney Ni and H

- (b) Nucleophilic substitution
- (d) Electrophilic substitution
- (b) Nascent hydrogen
- (d) Lithium hydride in ether

 CH_3

OH

229. What are the products obtained on hydroboration-oxidation of the given alkene

 CH_3

OH

 CH_3

HŌ

H

H

OH

 CH_3

(I)
$$CH_3$$
 (II) CH_3 (VI)

ÔН

(a) I and III

H

- (b) II and IV
- (c) II and VI

(III)

(d) III and V

ANSWERS — LEVEL 1															" AO
1.	(c)	2.	(d)	3.	(c)	4.	(d)	5.	(b)	6.	(b)	7.	(c)	8.	(c)
9.	(c)	10.	(d)	11.	(b)	12.	(d)	13.	(c)	14.	(b)	15.	(b)	16.	(c)
17.	(d)	18.	(d)	19.	(b)	20.	(d)	21.	(b)	22.	(a)	23.	(b)	24.	(b)
25.	(b)	26.	(b)	27.	(d)	28.	(b)	29.	(d)	30.	(b)	31.	(c)	32.	(b)
33.	(a)	34.	(b)	35.	(b)	36.	(b)	37.	(b)	38.	(b)	39.	(b)	40.	(b)
41.	(d)	42.	(e)	43.	(c)	44.	(c)	45.	(a)	46.	(c)	47.	(c)	48.	(b)
49.	(b)	50.	(b)	51.	(b)	52.	(a)	53.	(b)	54.	(d)	55.	(b)	56.	(c)
57.	(c)	58.	(b)	59.	(c)	60.	(a)	61.	(b)	62.	(d)	63.	(a)	64.	(b)
65.	(d)	66.	(b)	67.	(d)	68.	(a)	69.	(c)	70.	(d)	71.	(d)	72.	(c)
73.	(d)	74.	(c)	75.	(a)	76.	(c)	77.	(d)	78.	(b)	79.	(d)	80.	(d)
81.	(c)	82.	(b)	83.	(a)	84.	(a)	85.	(a)	86.	(d)	87.	(b)	88.	(b)
89.	(c)	90.	(d)	91.	(b)	92.	(b)	93.	(a)	94.	(a)	95.	(a)	96.	(d)
97.	(b)	98.	(a)	99.	(c)	100.	(d)	101.	(b)	102.	(b)	103.	(d)	104.	(b)
105.	(b)	106.	(c)	107.	(b)	108.	(b)	109.	(d)	110.	(d)	111.	(c)	112.	(b)
113.	(a)	114.	(b)	115.	(c)	116.	(a)	117.	(b)	118.	(b)	119.	(b)	120.	(d)
121.	(b)	122.	(c)	123.	(a)	124.	(b)	125.	(d)	126.	(a)	127.	(b)	128.	(c)
129.	(d)	130.	(a)	131.	(c)	132.	(d)	133.	(c)	134.	(a)	135.	(d)	136.	(c)
137.	(b)	138.	(d)	139.	(d)	140.	(b)	141.	(b)	142.	(b)	143.	(b)	144.	(b)
145.	(b)	146.	(c)	147.	(b)	148.	(a)	149.	(d)	150.	(b)	151.	(a)	152.	(d)
153.	(c)	154.	(c)	155.	(b)	156.	(c)	157.	(a)	158.	(a)	159.	(b)	160.	(c)
161.	(b)	162.	(c)	163	(b)	164.	(d)	165.	(d)	166.	(b)	167.	(b)	168.	(a)
169.	(c)	170.	(b)	171.	(b)	172.	(d)	173.	(b)	174.	(c)	175.	(b)	176.	(b)
177.	(b)	178.	(c)	179.	(b)	180.	(c)	181.	(b)	182.	(b)	183.	(c)	184.	(c)
185.	(c)	186.	(p)	187.	(a)	188.	(b)	189.	(b)	190.	(b)	191.	(a)	192.	(b)
193.	(c)	194.	(b)	195.	(c)	196.	(c)	197.	(b)	198.	(a)	199.	(b)	200.	
201.	(c)	202.	(c)	203.	(b)	204.	(d)	205.	(b)	206.		207.		208.	(b)
209.	(b)	210.	(b)	211.	(c)	212.	(c)	213.	(b)	214.	(a)	215.		216.	(b)
217.	(d)	218.	(c)	219.	(a)	220.	(b)	221.	(b)	222.		223.	_	224.	(c)
225.	(b,c)	226.	(b)	227.	(c)	228.	(c)	229.	(d)				1		17